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(54) Title: HUMAN RECEPTOR PROTEINS; RELATED REAGENTS AND METHODS

(57) Abstract: Nucleic acids encoding mammalian, e.g., human receptors, purified receptor proteins and fragments thereof. Antibodies, both polyclonal and monoclonal, are also provided. Methods of using the compositions for both diagnostic and therapeutic utilities are provided.

HUMAN RECEPTOR PROTEINS; RELATED REAGENTS AND METHODS

FIELD OF THE INVENTION

The present invention relates to compositions and methods for affecting mammalian physiology, including morphogenesis or immune system function. In particular, it provides nucleic acids, proteins, and antibodies which regulate development and/or the immune system. Diagnostic and therapeutic uses of these materials are also disclosed.

BACKGROUND OF THE INVENTION

Recombinant DNA technology refers generally to techniques of integrating genetic information from a donor source into vectors for subsequent processing, such as through introduction into a host, whereby the transferred genetic information is copied and/or expressed in the new environment. Commonly, the genetic information exists in the form of complementary DNA (cDNA) derived from messenger RNA (mRNA) coding for a desired protein product. The carrier is frequently a plasmid having the capacity to incorporate cDNA for later replication in a host and, in some cases, actually to control expression of the cDNA and thereby direct synthesis of the encoded product in the host.

For some time, it has been known that the mammalian immune response is based on a series of complex cellular interactions, called the "immune network". Recent research has provided new insights into the inner workings of this network. While it remains clear that much of the immune response does, in fact, revolve around the network-like interactions of lymphocytes, macrophages, granulocytes, and other cells, immunologists now generally hold the opinion that soluble proteins, known as lymphokines, cytokines, or monokines, play critical roles in controlling these cellular interactions. Thus, there is considerable interest in the isolation, characterization, and mechanisms of action of cell modulatory factors, an understanding of which will lead to significant advancements in the diagnosis and therapy of numerous medical abnormalities, e.g., immune system disorders.

Lymphokines apparently mediate cellular activities in a variety of ways. They have been shown to support the proliferation, growth, and/or differentiation of pluripotential hematopoietic stem cells into vast numbers of progenitors comprising diverse cellular lineages which make up a complex immune system. Proper and balanced interactions between the cellular components are necessary for a healthy immune response. The different cellular lineages often respond in a different manner when lymphokines are administered in conjunction with other agents.

Cell lineages especially important to the immune response include two classes of lymphocytes: B-cells, which can produce and secrete immunoglobulins (proteins with the capability of recognizing and binding to foreign matter to effect its removal), and T-cells of various subsets that secrete lymphokines and induce or suppress the B-cells and various other cells (including other T-

cells) making up the immune network. These lymphocytes interact with many other cell types.

Another important cell lineage is the mast cell (which has not been positively identified in all mammalian species), which is a granule-containing connective tissue cell located proximal to capillaries throughout the body. These cells are found in especially high concentrations in the lungs, skin, and gastrointestinal and genitourinary tracts. Mast cells play a central role in allergy-related disorders, particularly anaphylaxis as follows: when selected antigens crosslink one class of immunoglobulins bound to receptors on the mast cell surface, the mast cell degranulates and releases mediators, e.g., histamine, serotonin, heparin, and prostaglandins, which cause allergic reactions, e.g., anaphylaxis.

Research to better understand and treat various immune disorders has been hampered by the general inability to maintain cells of the immune system in vitro. Immunologists have discovered that culturing many of these cells can be accomplished through the use of T-cell and other cell supernatants, which contain various growth factors, including many of the lymphokines.

The interleukin-1 family of proteins includes the IL-1α, the IL-1β, the IL-1RA, and recently the IL-1γ (also designated Interferon-Gamma Inducing Factor, IGIF). This related family of genes have been implicated in a broad range of biological functions. See Dinarello (1994) FASEB J. 8:1314-1325; Dinarello (1991) Blood 77:1627-1652; and Okamura, et al. (1995) Nature 378:88-91.

In addition, various growth and regulatory factors exist which modulate morphogenetic development. This includes, e.g., the Toll ligands, which signal through binding to receptors which share structural, and mechanistic, features characteristic of the IL-1 receptors. See, e.g., Lemaitre, et al. (1996) Cell

86:973-983; and Belvin and Anderson (1996) Ann. Rev. Cell & Devel. Biol. 12:393-416.

From the foregoing, it is evident that the discovery and development of new soluble proteins and their receptors, including ones similar to lymphokines, should contribute to new therapies for a wide range of degenerative or abnormal conditions which directly or indirectly involve development, differentiation, or function, e.g., of the immune system and/or hematopoietic cells. In particular, the discovery and understanding of novel receptors for lymphokine-like molecules which enhance or potentiate the beneficial activities of other lymphokines would be highly advantageous. The present invention provides new receptors for ligands exhibiting similarity to interleukin-1 like compositions and related compounds, and methods for their use.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 shows a schematic comparison of the protein architectures of Drosophila, Caenorabditis, and human DTLRs, and their relationship to vertebrate IL-1 receptors and plant disease resistance proteins. Three Drosophila (Dm) DTLRs (Toll, 18w, and the Mst ORF fragment) (Morisato and Anderson (1995) Ann. Rev. Genet. 29:371-399; Chiang and Beachy (1994) Mech. Develop. 47:225-239; Mitcham, et al. (1996) J. Biol. Chem. 271:5777-5783; and Eldon, et al. (1994) Develop. 120:885-899) are arrayed beside four complete (DTLRs 1-4) and one partial (DTLR5) human (Hu) receptors. Individual LRRs in the receptor ectodomains that are flagged by PRINTS (Attwood, et al. (1997) Nucleic Acids Res. 25:212-217) are explicitly noted by boxes; 'top' and 'bottom' Cys-rich clusters that flank the C- or N-terminal ends of LRR arrays are respectively drawn by opposed half-circles. The loss of the internal Cys-rich region in DTLRs 1-5 largely accounts for their smaller ectodomains (558, 570, 690, and 652 aa, respectively) when compared to the 784 and 977 aa extensions of Toll and 18w. The incomplete chains of DmMst and HuDTLR5 (about 519 and 153 aa ectodomains, respectively) are represented by dashed lines. The intracellular signaling module common to DTLRs, IL-1-type receptors (IL-1Rs), the intracellular protein Myd88, and the tobacco disease resistance gene N product (DRgN) is indicated below the membrane. See, e.g., Hardiman, et al. (1996) Oncogene 13:2467-2475; and Rock, et al. (1998) Proc. Nat'l Acad. Sci. USA 95:588-. Additional domains include the trio of Ig-like modules in IL-1Rs (disulfide-linked loops); the DRgN protein features an NTPase domain (box) and Myd88 has a death domain (black oval).

Figures 2A-2C show conserved structural patterns in the signaling domains of Toll- and IL-1-like cytokine receptors, and two divergent modular proteins. Figures 2A-2B show a sequence alignment of the common TH domain.

DTLRs are labeled as in Figure 1; the human (Hu) or mouse (Mo) IL-1 family receptors (IL-1R1-6) are sequentially numbered as earlier proposed (Hardiman, et al. (1996) Oncogene 13:2467-2475); Myd88 and the sequences from tobacco (To) and flax, L. usitatissimum (Lu), represent C- and N-terminal domains, respectively, of larger, multidomain molecules. Ungapped blocks of sequence (numbered 1-10) are boxed. Triangles indicate deleterious mutations, while truncations N-terminal of the arrow eliminate bioactivity in human IL-1R1 (Heguy, et al. (1992) J. Biol. Chem. 267:2605-2609). PHD (Rost and Sander (1994) Proteins 19:55-72) and DSC (King and Sternberg (1996) Protein Sci. 5:2298-2310) secondary structure predictions of α-helix (H), β-strand (E), or coil (L) are marked. The amino acid shading scheme depicts chemically similar residues: hydrophobic, acidic, basic, Cys, aromatic, structure-breaking, and tiny. Diagnostic sequence patterns for IL-1Rs, DTLRs, and full alignment (ALL) were derived by Consensus at a stringency of 75%. Symbols for amino acid subsets are (see internet site for detail): o, alcohol; l, aliphatic; •, any amino acid; a, aromatic; c, charged; h, hydrophobic; -, negative; p, polar; +, positive; s, small; u, tiny; t, turnlike. Figure 2C shows a topology diagram of the proposed TH β/α domain fold. The parallel β-sheet (with β-strands A-E as yellow triangles) is seen at its C-terminal end; α-helices (circles labeled 1-5) link the β-strands; chain connections are to the front (visible) or back (hidden). Conserved, charged residues at the C-end of the β-sheet are noted in gray (Asp) or as a lone black (Arg) residue (see text).

Figure 3 shows evolution of a signaling domain superfamily. The multiple TH module alignment of Figures 2A-2B was used to derive a phylogenetic tree by the Neighbor-Joining method (Thompson, et al. (1994) Nucleic

Acids Res. 22:4673-4680). Proteins labeled as in the alignment; the tree was rendered with TreeView.

Figures 4A-4D depict FISH chromosomal mapping of human DTLR genes. Denatured chromosomes from synchronous cultures of human lymphocytes were hybridized to biotinylated DTLR cDNA probes for localization. The assignment of the FISH mapping data (left, Figures 4A, DTLR2; 4B, DTLR3; 4C, DTLR4; 4D, DTLR5) with chromosomal bands was achieved by superimposing FISH signals with DAPI banded chromosomes (center panels). Heng and Tsui (1994) Meth. Molec. Biol. 33:109-122. Analyses are summarized in the form of human chromosome ideograms (right panels).

Figures 5A-5F depict mRNA blot analyses of Human DTLRs. Human multiple tissue blots (He, heart; Br, brain; Pl, placenta; Lu, lung; Li, liver; Mu, muscle; Ki, kidney; Pn, Pancreas; Sp, spleen; Th, thymus; Pr, prostate; Te, testis; Ov, ovary, SI, small intestine; Co, colon; PBL, peripheral blood lymphocytes) and cancer cell line (promyelocytic leukemia, HL60; cervical cancer, HELAS3; chronic myelogenous leukemia, K562; lymphoblastic leukemia, Molt4; colorectal adenocarcinoma, SW480; melanoma, G361; Burkitt's Lymphoma Raji, Burkitt's; colorectal adenocarcinoma, SW480; lung carcinoma, A549) containing approximately 2 µg of poly(A)$^+$ RNA per lane were probed with radiolabeled cDNAs encoding DTLR1 (Figures 5A-5C), DTLR2 (Figure 5D), DTLR3 (Figure 5E), and DTLR4 (Figure 5F) as described. Blots were exposed to X-ray film for 2 days (Figures 5A-5C) or one week (Figure 5D-5F) at -70° C with intensifying screens. An anomalous 0.3 kB species appears in some lanes; hybridization experiments exclude a message encoding a DTLR cytoplasmic fragment.

SUMMARY OF THE INVENTION

The present invention is directed to nine novel related mammalian receptors, e.g., primate, human, DNAX Toll receptor like molecular structures, designated DTLR2, DTLR3, DTLR4, DTLR5, DTLR7, DTLR8, DTLR9, and DTLR10, and their biological activities. It includes nucleic acids coding for the polypeptides themselves and methods for their production and use. The nucleic acids of the invention are characterized, in part, by their homology to cloned complementary DNA (cDNA) sequences enclosed herein.

In certain embodiments, the invention provides a composition of matter selected from the group of: a substantially pure or recombinant DTLR2 protein or peptide exhibiting identity over a length of at least about 12 amino acids to SEQ ID NO: 4; a natural sequence DTLR2 of SEQ ID NO: 4; a fusion protein comprising DTLR2 sequence; a substantially pure or recombinant DTLR3 protein or peptide exhibiting identity over a length of at least about 12 amino acids to SEQ ID NO: 6; a natural sequence DTLR3 of SEQ ID NO: 6; a fusion protein comprising DTLR3 sequence; a substantially pure or recombinant DTLR4 protein or peptide exhibiting identity over a length of at least about 12 amino acids to SEQ ID NO: 26; a natural sequence DTLR4 of SEQ ID NO: 26; a fusion protein comprising DTLR4 sequence; a substantially pure or recombinant DTLR5 protein or peptide exhibiting identity over a length of at least about 12 amino acids to SEQ ID NO: 10; a natural sequence DTLR5 of SEQ ID NO: 10; a fusion protein comprising DTLR5 sequence; a substantially pure or recombinant DTLR6 protein or peptide exhibiting identity over a length of at least about 12 amino acids to SEQ ID NO: 12, 28, or 30; a natural sequence DTLR6 of SEQ ID NO: 12, 28, or 30; a fusion protein comprising DTLR6 sequence; a substantially pure or recombinant DTLR7

protein or peptide exhibiting identity over a length of at least about 12 amino acids to SEQ ID NO: 16, 18, or 37; a natural sequence DTLR7 of SEQ ID NO: 16, 18, or 37; a fusion protein comprising DTLR7 sequence; a substantially pure or recombinant DTLR8 protein or peptide exhibiting identity over a length of at least about 12 amino acids to SEQ ID NO: 32 or 39; a natural sequence DTLR8 of SEQ ID NO: 32 or 39; a fusion protein comprising DTLR8 sequence; a substantially pure or recombinant DTLR9 protein or peptide exhibiting identity over a length of at least about 12 amino acids to SEQ ID NO: 22 or 41; a natural sequence DTLR9 of SEQ ID NO: 22 or 41; a fusion protein comprising DTLR9 sequence; a substantially pure or recombinant DTLR10 protein or peptide exhibiting identity over a length of at least about 12 amino acids to SEQ ID NO: 34, 43, or 45; a natural sequence DTLR10 of SEQ ID NO: 34, 43, or 45; and a fusion protein comprising DTLR10 sequence. Preferably, the substantially pure or isolated protein comprises a segment exhibiting sequence identity to a corresponding portion of a DTLR2, DTLR3, DTLR4, DTLR5, DTLR6, DTLR7, DTLR8, DTLR9, or DTLR10, wherein said identity is over at least about 15 amino acids; preferably about 19 amino acids; or more preferably about 25 amino acids. In specific embodiments, the composition of matter: is DTLR2, which comprises a mature sequence of Table 2; or lacks a post-translational modification; is DTLR3, which comprises a mature sequence of Table 3; or lacks a post-translational modification; is DTLR4, which: comprises a mature sequence of Table 4; or lacks a post-translational modification; is DTLR5, which: comprises the complete sequence of Table 5; or lacks a post-translational; is DTLR6, which comprises a mature sequence of Table 6; or lacks a post-translational modification; is DTLR7, which comprises a mature sequence of Table 7; or lacks a post-translational modification; is DTLR8, which: comprises a mature sequence of Table 8; or lacks a post-

translational modification; is DTLR9, which: comprises the complete sequence of Table 9; or lacks a post-translational; is DTLR10, which comprises a mature sequence of Table 10; or lacks a post-translational modification; or the composition of matter may be a protein or peptide which: is from a warm blooded animal selected from a mammal, including a primate, such as a human; comprises at least one polypeptide segment of SEQ ID NO: 4, 6, 26, 10, 12, 28, 30, 16, 18, 32, 22, or 34; exhibits a plurality of portions exhibiting said identity; is a natural allelic variant of DTLR2, DTLR3, DTLR4, DTLR5, DTLR6, DTLR7, DTLR8, DTLR9, or DTLR10; has a length at least about 30 amino acids; exhibits at least two non-overlapping epitopes which are specific for a primate DTLR2, DTLR3, DTLR4, DTLR5, DTLR6, DTLR7, DTLR8, DTLR9, or DTLR10; exhibits sequence identity over a length of at least about 35 amino acids to a primate DTLR2, DTLR3, DTLR4, DTLR5, DTLR6, DTLR7, DTLR8, DTLR9. or DTLR10; further exhibits at least two non-overlapping epitopes which are specific for a primate DTLR2, DTLR3, DTLR4, DTLR5, DTLR6, DTLR7, DTLR8, DTLR9, or DTLR10; exhibits identity over a length of at least about 20 amino acids to a rodent DTLR6; is glycosylated; has a molecular weight of at least 100 kD with natural glycosylation; is a synthetic polypeptide; is attached to a solid substrate; is conjugated to another chemical moiety; is a 5-fold or less substitution from natural sequence; or is a deletion or insertion variant from a natural sequence.

Other embodiments include a composition comprising: a sterile DTLR2 protein or peptide; or the DTLR2 protein or peptide and a carrier, wherein the carrier is: an aqueous compound, including water, saline, and/or buffer; and/or formulated for oral, rectal, nasal, topical, or parenteral administration; a sterile DTLR3 protein or peptide; or the DTLR3 protein or peptide and a carrier, wherein the carrier is: an aqueous compound, including water, saline,

and/or buffer; and/or formulated for oral, rectal, nasal, topical, or parenteral administration; a sterile DTLR4 protein or peptide; or the DTLR4 protein or peptide and a carrier, wherein the carrier is: an aqueous compound, including water, saline, and/or buffer; and/or formulated for oral, rectal, nasal, topical, or parenteral administration; a sterile DTLR5 protein or peptide; or the DTLR5 protein or peptide and a carrier, wherein the carrier is: an aqueous compound, including water, saline, and/or buffer; and/or formulated for oral, rectal, nasal, topical, or parenteral administration; a sterile DTLR6 protein or peptide; or the DTLR6 protein or peptide and a carrier, wherein the carrier is: an aqueous compound, including water, saline, and/or buffer; and/or formulated for oral, rectal, nasal, topical, or parenteral administration; a sterile DTLR7 protein or peptide; or the DTLR7 protein or peptide and a carrier, wherein the carrier is: an aqueous compound, including water, saline, and/or buffer; and/or formulated for oral, rectal, nasal, topical, or parenteral administration; a sterile DTLR8 protein or peptide; or the DTLR8 protein or peptide and a carrier, wherein the carrier is: an aqueous compound, including water, saline, and/or buffer; and/or formulated for oral, rectal, nasal, topical, or parenteral administration; a sterile DTLR9 protein or peptide; or the DTLR9 protein or peptide and a carrier, wherein the carrier is: an aqueous compound, including water, saline, and/or buffer; and/or formulated for oral, rectal, nasal, topical, or parenteral administration; a sterile DTLR10 protein or peptide; or the DTLR10 protein or peptide and a carrier, wherein the carrier is: an aqueous compound, including water, saline, and/or buffer; and/or formulated for oral, rectal, nasal, topical, or parenteral administration.

In certain fusion protein embodiments, the invention provides a fusion protein comprising: mature protein

sequence of Table 2, 3, 4, 5, 6, 7, 8, 9, or 10; a detection or purification tag, including a FLAG, His6, or Ig sequence; or sequence of another receptor protein.

Various kit embodiments include a kit comprising a DTLR protein or polypeptide, and: a compartment comprising the protein or polypeptide; and/or instructions for use or disposal of reagents in the kit.

Binding compound embodiments include those comprising an antigen binding site from an antibody, which specifically binds to a natural DTLR2, DTLR3, DTLR4, DTLR5, DTLR6, DTLR7, DTLR8, DTLR9, or DTLR10 protein, wherein: the protein is a primate protein; the binding compound is an Fv, Fab, or Fab2 fragment; the binding compound is conjugated to another chemical moiety; or the antibody: is raised against a peptide sequence of a mature polypeptide of Table 2, 3, 4, 5, 6, 7, 8, 9, or 10; is raised against a mature DTLR2, DTLR3, DTLR4, DTLR5, DTLR6, DTLR7, DTLR8, DTLR9, or DTLR10; is raised to a purified human DTLR2, DTLR3, DTLR4, DTLR5, DTLR6, DTLR7, DTLR8, DTLR9, or DTLR10; is immunoselected; is a polyclonal antibody; binds to a denatured DTLR2, DTLR3, DTLR4, DTLR5, DTLR6, DTLR7, DTLR8, DTLR9, or DTLR10; exhibits a Kd to antigen of at least 30 μM; is attached to a solid substrate, including a bead or plastic membrane; is in a sterile composition; or is detectably labeled, including a radioactive or fluorescent label. A binding composition kit often comprises the binding compound, and: a compartment comprising said binding compound; and/or instructions for use or disposal of reagents in the kit. Often the kit is capable of making a qualitative or quantitative analysis.

Methods are provided, e.g., of making an antibody, comprising immunizing an immune system with an immunogenic amount of a primate DTLR2, DTLR3, DTLR4, DTLR5, DTLR6, DTLR7, DTLR8, DTLR9, or DTLR10, thereby causing said antibody to be produced; or producing an antigen:antibody

complex, comprising contacting such an antibody with a mammalian DTLR2, DTLR3, DTLR4, DTLR5, DTLR6, DTLR7, DTLR8, DTLR9, or DTLR10 protein or peptide, thereby allowing said complex to form.

Other compositions include a composition comprising: a sterile binding compound, or the binding compound and a carrier, wherein the carrier is: an aqueous compound, including water, saline, and/or buffer; and/or formulated for oral, rectal, nasal, topical, or parenteral administration.

Nucleic acid embodiments include an isolated or recombinant nucleic acid encoding a DTLR2-10 protein or peptide or fusion protein, wherein: the DTLR is from a mammal; or the nucleic acid: encodes an antigenic peptide sequence of Table 2, 3, 4, 5, 6, 7, 8, 9, or 10; encodes a plurality of antigenic peptide sequences of Table 2, 3, 4, 5, 6, 7, 8, 9, or 10; comprises at least 17 contiguous nucleotides from Table 2, 3, 4, 5, 6, 7, 8, 9, or 10; exhibits at least about 80% identity to a natural cDNA encoding said segment; is an expression vector; further comprises an origin of replication; is from a natural source; comprises a detectable label; comprises synthetic nucleotide sequence; is less than 6 kb, preferably less than 3 kb; is from a mammal, including a primate; comprises a natural full length coding sequence; is a hybridization probe for a gene encoding said DTLR; or is a PCR primer, PCR product, or mutagenesis primer. A cell, tissue, or organ comprising such a recombinant nucleic acid is also provided. Preferably, the cell is: a prokaryotic cell; a eukaryotic cell; a bacterial cell; a yeast cell; an insect cell; a mammalian cell; a mouse cell; a primate cell; or a human cell. Kits are provided comprising such nucleic acids, and: a compartment comprising said nucleic acid; a compartment further comprising a primate DTLR2, DTLR3, DTLR4, or DTLR5 protein or polypeptide; and/or instructions for use or disposal of

reagents in the kit. Often, the kit is capable of making a qualitative or quantitative analysis.

Other embodiments include a nucleic acid which: hybridizes under wash conditions of 30° C and less than 2M salt to SEQ ID NO: 3; hybridizes under wash conditions of 30° C and less than 2 M salt to SEQ ID NO: 5; hybridizes under wash conditions of 30° C and less than 2M salt to SEQ ID NO: 7; hybridizes under wash conditions of 30° C and less than 2 M salt to SEQ ID NO: 9; hybridizes under wash conditions of 30° C and less than 2 M salt to SEQ ID NO: 11, 13, 27, or 29; hybridizes under wash conditions of 30° C and less than 2 M salt to SEQ ID NO: 15, 17, or 36; hybridizes under wash conditions of 30° C and less than 2 M salt to SEQ ID NO: 19, 31, or 38; hybridizes under wash conditions of 30° C and less than 2 M salt to SEQ ID NO: 21 or 40; hybridizes under wash conditions of 30° C and less than 2 M salt to SEQ ID NO: 23, 33, 42, or 44; exhibits at least about 85% identity over a stretch of at least about 30 nucleotides to a primate DTLR2; exhibits at least about 85% identity over a stretch of at least about 30 nucleotides to a primate DTLR3; exhibits at least about 85% identity over a stretch of at least about 30 nucleotides to a primate DTLR4; or exhibits at least about 85% identity over a stretch of at least about 30 nucleotides to a primate DTLR5. Preferably, such nucleic acid will have such properties, wherein: wash conditions are at 45° C and/or 500 mM salt; or the identity is at least 90% and/or the stretch is at least 55 nucleotides.

More preferably, the wash conditions are at 55° C and/or 150 mM salt; or the identity is at least 95% and/or the stretch is at least 75 nucleotides.

Also provided are methods of producing a ligand:receptor complex, comprising contacting a substantially pure primate DTLR2, DTLR3, DTLR4, DTLR5, DTLR6, DTLR7, DTLR8, DTLR9, or DTLR10, including a recombinant or synthetically produced protein, with

candidate Toll ligand; thereby allowing said complex to form.

The invention also provides a method of modulating physiology or development of a cell or tissue culture cells comprising contacting the cell with an agonist or antagonist of a mammalian DTLR2, DTLR3, DTLR4, DTLR5, DTLR6, DTLR7, DTLR8, DTLR9, or DTLR10. Preferably, the cell is a pDC2 cell with the agonist or antagonist of DTLR10.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

OUTLINE

I. General
II. Activities
III. Nucleic acids
  A. encoding fragments, sequence, probes
  B. mutations, chimeras, fusions
  C. making nucleic acids
  D. vectors, cells comprising
IV. Proteins, Peptides
  A. fragments, sequence, immunogens, antigens
  B. muteins
  C. agonists/antagonists, functional equivalents
  D. making proteins
V. Making nucleic acids, proteins
  A. synthetic
  B. recombinant
  C. natural sources
VI. Antibodies
  A. polyclonals
  B. monoclonal
  C. fragments; Kd
  D. anti-idiotypic antibodies
  E. hybridoma cell lines
VII. Kits and Methods to quantify DTLRs 2-10
  A. ELISA
  B. assay mRNA encoding
  C. qualitative/quantitative
  D. kits
VIII. Therapeutic compositions, methods
  A. combination compositions
  B. unit dose
  C. administration
IX. Ligands

I. General

The present invention provides the amino acid sequence and DNA sequence of mammalian, herein primate DNAX Toll like receptor molecules (DTLR) having particular defined properties, both structural and biological. These have been designated herein as DTLR2, DTLR3, DTLR4, DTLR5, DTLR6, DTLR7, DTLR8, DTLR9, and DTLR10, respectively, and increase the number of members of the human Toll like

receptor family from 1 to 10. Various cDNAs encoding these molecules were obtained from primate, e.g., human, cDNA sequence libraries. Other primate or other mammalian counterparts would also be desired.

Some of the standard methods applicable are described or referenced, e.g., in Maniatis, et al. (1982) Molecular Cloning, A Laboratory Manual, Cold Spring Harbor Laboratory, Cold Spring Harbor Press; Sambrook, et al. (1989) Molecular Cloning: A Laboratory Manual, (2d ed.), vols. 1-3, CSH Press, NY; Ausubel, et al., Biology, Greene Publishing Associates, Brooklyn, NY; or Ausubel, et al. (1987 and periodic supplements) Current Protocols in Molecular Biology, Greene/Wiley, New York; each of which is incorporated herein by reference.

A complete nucleotide (SEQ ID NO: 1) and corresponding amino acid sequence (SEQ ID NO: 2) of a human DTLR1 coding segment is shown in Table 1. See also Nomura, et al. (1994) DNA Res. 1:27-35. A complete nucleotide (SEQ ID NO: 3) and corresponding amino acid sequence (SEQ ID NO: 4) of a human DTLR2 coding segment is shown in Table 2. A complete nucleotide (SEQ ID NO: 5) and corresponding amino acid sequence (SEQ ID NO: 6) of a human DTLR3 coding segment is shown in Table 3. A complete nucleotide (SEQ ID NO: 7) and corresponding amino acid sequence (SEQ ID NO: 8) of a human DTLR4 coding segment is shown in Table 4; see also SEQ ID NO: 25 and 26. A partial nucleotide (SEQ ID NO: 9) and corresponding amino acid sequence (SEQ ID NO: 10) of a human DTLR5 coding segment is shown in Table 5. A complete nucleotide (SEQ ID NO: 11) and corresponding amino acid sequence (SEQ ID NO: 12) of a human DTLR6 coding segment is shown in Table 6, along with partial sequence of a mouse DTLR6 (SEQ ID NO: 13, 14, 27, 28, 29, and 30). Partial nucleotide (SEQ ID NO: 15 and 17) and corresponding amino acid sequence (SEQ ID NO: 16 and 18) of a human DTLR7 coding segment is shown in Table 7; full length sequence is

provided in SEQ ID NO: 36 and 37. Partial nucleotide (SEQ ID NO: 19) and corresponding amino acid sequence (SEQ ID NO: 20) of a human DTLR8 coding segment is shown in Table 8, with supplementary sequence (SEQ ID NO: 31, 32, 38, and 39). Partial nucleotide (SEQ ID NO: 21) and corresponding amino acid sequence (SEQ ID NO: 22) of a human DTLR9 coding segment is shown in Table 9; see also SEQ ID NO: 40 and 41. Partial nucleotide (SEQ ID NO: 23) and corresponding amino acid sequence (SEQ ID NO: 24) of a human DTLR10 coding segment is shown in Table 10, along with supplementary sequence (SEQ ID NO: 33, 34, 42, and 43) and rodent, e.g., mouse, sequence (SEQ ID NO: 35, 44, and 45).

Table 1: Nucleotide and amino acid sequences (see SEQ ID NO: 1 and 2) of a primate, e.g., human, DNAX Toll like receptor 1 (DTLR1).

```
ATG ACT AGC ATC TTC CAT TTT GCC ATT ATC TTC ATG TTA ATA CTT CAG        48
Met Thr Ser Ile Phe His Phe Ala Ile Ile Phe Met Leu Ile Leu Gln
-22  -20                -15                 -10

ATC AGA ATA CAA TTA TCT GAA GAA AGT GAA TTT TTA GTT GAT AGG TCA        96
Ile Arg Ile Gln Leu Ser Glu Glu Ser Glu Phe Leu Val Asp Arg Ser
 -5                   1                   5                  10

AAA AAC GGT CTC ATC CAC GTT CCT AAA GAC CTA TCC CAG AAA ACA ACA       144
Lys Asn Gly Leu Ile His Val Pro Lys Asp Leu Ser Gln Lys Thr Thr
                15                  20                  25

ATC TTA AAT ATA TCG CAA AAT TAT ATA TCT GAG CTT TGG ACT TCT GAC       192
Ile Leu Asn Ile Ser Gln Asn Tyr Ile Ser Glu Leu Trp Thr Ser Asp
            30                  35                  40

ATC TTA TCA CTG TCA AAA CTG AGG ATT TTG ATA ATT TCT CAT AAT AGA       240
Ile Leu Ser Leu Ser Lys Leu Arg Ile Leu Ile Ile Ser His Asn Arg
        45                  50                  55

ATC CAG TAT CTT GAT ATC AGT GTT TTC AAA TTC AAC CAG GAA TTG GAA       288
Ile Gln Tyr Leu Asp Ile Ser Val Phe Lys Phe Asn Gln Glu Leu Glu
    60                  65                  70

TAC TTG GAT TTG TCC CAC AAC AAG TTG GTG AAG ATT TCT TGC CAC CCT       336
Tyr Leu Asp Leu Ser His Asn Lys Leu Val Lys Ile Ser Cys His Pro
75                  80                  85                  90

ACT GTG AAC CTC AAG CAC TTG GAC CTG TCA TTT AAT GCA TTT GAT GCC       384
Thr Val Asn Leu Lys His Leu Asp Leu Ser Phe Asn Ala Phe Asp Ala
                95                 100                 105

CTG CCT ATA TGC AAA GAG TTT GGC AAT ATG TCT CAA CTA AAA TTT CTG       432
Leu Pro Ile Cys Lys Glu Phe Gly Asn Met Ser Gln Leu Lys Phe Leu
            110                 115                 120

GGG TTG AGC ACC ACA CAC TTA GAA AAA TCT AGT GTG CTG CCA ATT GCT       480
Gly Leu Ser Thr Thr His Leu Glu Lys Ser Ser Val Leu Pro Ile Ala
        125                 130                 135

CAT TTG AAT ATC AGC AAG GTC TTG CTG GTC TTA GGA GAG ACT TAT GGG       528
His Leu Asn Ile Ser Lys Val Leu Leu Val Leu Gly Glu Thr Tyr Gly
    140                 145                 150

GAA AAA GAA GAC CCT GAG GGC CTT CAA GAC TTT AAC ACT GAG AGT CTG       576
Glu Lys Glu Asp Pro Glu Gly Leu Gln Asp Phe Asn Thr Glu Ser Leu
155                 160                 165                 170

CAC ATT GTG TTC CCC ACA AAC AAA GAA TTC CAT TTT ATT TTG GAT GTG       624
His Ile Val Phe Pro Thr Asn Lys Glu Phe His Phe Ile Leu Asp Val
                175                 180                 185
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TCA GTC AAG ACT GTA GCA AAT CTG GAA CTA TCT AAT ATC AAA TGT GTG      672
Ser Val Lys Thr Val Ala Asn Leu Glu Leu Ser Asn Ile Lys Cys Val
            190                 195                 200

CTA GAA GAT AAC AAA TGT TCT TAC TTC CTA AGT ATT CTG GCG AAA CTT      720
Leu Glu Asp Asn Lys Cys Ser Tyr Phe Leu Ser Ile Leu Ala Lys Leu
        205                 210                 215

CAA ACA AAT CCA AAG TTA TCA AGT CTT ACC TTA AAC AAC ATT GAA ACA      768
Gln Thr Asn Pro Lys Leu Ser Ser Leu Thr Leu Asn Asn Ile Glu Thr
    220                 225                 230

ACT TGG AAT TCT TTC ATT AGG ATC CTC CAA CTA GTT TGG CAT ACA ACT      816
Thr Trp Asn Ser Phe Ile Arg Ile Leu Gln Leu Val Trp His Thr Thr
235                 240                 245                 250

GTA TGG TAT TTC TCA ATT TCA AAC GTG AAG CTA CAG GGT CAG CTG GAC      864
Val Trp Tyr Phe Ser Ile Ser Asn Val Lys Leu Gln Gly Gln Leu Asp
                255                 260                 265

TTC AGA GAT TTT GAT TAT TCT GGC ACT TCC TTG AAG GCC TTG TCT ATA      912
Phe Arg Asp Phe Asp Tyr Ser Gly Thr Ser Leu Lys Ala Leu Ser Ile
            270                 275                 280

CAC CAA GTT GTC AGC GAT GTG TTC GGT TTT CCG CAA AGT TAT ATC TAT      960
His Gln Val Val Ser Asp Val Phe Gly Phe Pro Gln Ser Tyr Ile Tyr
        285                 290                 295

GAA ATC TTT TCG AAT ATG AAC ATC AAA AAT TTC ACA GTG TCT GGT ACA     1008
Glu Ile Phe Ser Asn Met Asn Ile Lys Asn Phe Thr Val Ser Gly Thr
    300                 305                 310

CGC ATG GTC CAC ATG CTT TGC CCA TCC AAA ATT AGC CCG TTC CTG CAT     1056
Arg Met Val His Met Leu Cys Pro Ser Lys Ile Ser Pro Phe Leu His
315                 320                 325                 330

TTG GAT TTT TCC AAT AAT CTC TTA ACA GAC ACG GTT TTT GAA AAT TGT     1104
Leu Asp Phe Ser Asn Asn Leu Leu Thr Asp Thr Val Phe Glu Asn Cys
                335                 340                 345

GGG CAC CTT ACT GAG TTG GAG ACA CTT ATT TTA CAA ATG AAT CAA TTA     1152
Gly His Leu Thr Glu Leu Glu Thr Leu Ile Leu Gln Met Asn Gln Leu
            350                 355                 360

AAA GAA CTT TCA AAA ATA GCT GAA ATG ACT ACA CAG ATG AAG TCT CTG     1200
Lys Glu Leu Ser Lys Ile Ala Glu Met Thr Thr Gln Met Lys Ser Leu
        365                 370                 375

CAA CAA TTG GAT ATT AGC CAG AAT TCT GTA AGC TAT GAT GAA AAG AAA     1248
Gln Gln Leu Asp Ile Ser Gln Asn Ser Val Ser Tyr Asp Glu Lys Lys
    380                 385                 390

GGA GAC TGT TCT TGG ACT AAA AGT TTA TTA AGT TTA AAT ATG TCT TCA     1296
Gly Asp Cys Ser Trp Thr Lys Ser Leu Leu Ser Leu Asn Met Ser Ser
395                 400                 405                 410
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AAT ATA CTT ACT GAC ACT ATT TTC AGA TGT TTA CCT CCC AGG ATC AAG      1344
Asn Ile Leu Thr Asp Thr Ile Phe Arg Cys Leu Pro Pro Arg Ile Lys
                415                 420                 425

GTA CTT GAT CTT CAC AGC AAT AAA ATA AAG AGC ATT CCT AAA CAA GTC      1392
Val Leu Asp Leu His Ser Asn Lys Ile Lys Ser Ile Pro Lys Gln Val
            430                 435                 440

GTA AAA CTG GAA GCT TTG CAA GAA CTC AAT GTT GCT TTC AAT TCT TTA      1440
Val Lys Leu Glu Ala Leu Gln Glu Leu Asn Val Ala Phe Asn Ser Leu
        445                 450                 455

ACT GAC CTT CCT GGA TGT GGC AGC TTT AGC AGC CTT TCT GTA TTG ATC      1488
Thr Asp Leu Pro Gly Cys Gly Ser Phe Ser Ser Leu Ser Val Leu Ile
    460                 465                 470

ATT GAT CAC AAT TCA GTT TCC CAC CCA TCA GCT GAT TTC TTC CAG AGC      1536
Ile Asp His Asn Ser Val Ser His Pro Ser Ala Asp Phe Phe Gln Ser
475                 480                 485                 490

TGC CAG AAG ATG AGG TCA ATA AAA GCA GGG GAC AAT CCA TTC CAA TGT      1584
Cys Gln Lys Met Arg Ser Ile Lys Ala Gly Asp Asn Pro Phe Gln Cys
                495                 500                 505

ACC TGT GAG CTC GGA GAA TTT GTC AAA AAT ATA GAC CAA GTA TCA AGT      1632
Thr Cys Glu Leu Gly Glu Phe Val Lys Asn Ile Asp Gln Val Ser Ser
            510                 515                 520

GAA GTG TTA GAG GGC TGG CCT GAT TCT TAT AAG TGT GAC TAC CCG GAA      1680
Glu Val Leu Glu Gly Trp Pro Asp Ser Tyr Lys Cys Asp Tyr Pro Glu
        525                 530                 535

AGT TAT AGA GGA ACC CTA CTA AAG GAC TTT CAC ATG TCT GAA TTA TCC      1728
Ser Tyr Arg Gly Thr Leu Leu Lys Asp Phe His Met Ser Glu Leu Ser
    540                 545                 550

TGC AAC ATA ACT CTG CTG ATC GTC ACC ATC GTT GCC ACC ATG CTG GTG      1776
Cys Asn Ile Thr Leu Leu Ile Val Thr Ile Val Ala Thr Met Leu Val
555                 560                 565                 570

TTG GCT GTG ACT GTG ACC TCC CTC TGC ATC TAC TTG GAT CTG CCC TGG      1824
Leu Ala Val Thr Val Thr Ser Leu Cys Ile Tyr Leu Asp Leu Pro Trp
                575                 580                 585

TAT CTC AGG ATG GTG TGC CAG TGG ACC CAG ACC CGG CGC AGG GCC AGG      1872
Tyr Leu Arg Met Val Cys Gln Trp Thr Gln Thr Arg Arg Arg Ala Arg
            590                 595                 600

AAC ATA CCC TTA GAA GAA CTC CAA AGA AAT CTC CAG TTT CAT GCA TTT      1920
Asn Ile Pro Leu Glu Glu Leu Gln Arg Asn Leu Gln Phe His Ala Phe
        605                 610                 615
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ATT TCA TAT AGT GGG CAC GAT TCT TTC TGG GTG AAG AAT GAA TTA TTG      1968
Ile Ser Tyr Ser Gly His Asp Ser Phe Trp Val Lys Asn Glu Leu Leu
    620                 625                 630

CCA AAC CTA GAG AAA GAA GGT ATG CAG ATT TGC CTT CAT GAG AGA AAC      2016
Pro Asn Leu Glu Lys Glu Gly Met Gln Ile Cys Leu His Glu Arg Asn
635                 640                 645                 650

TTT GTT CCT GGC AAG AGC ATT GTG GAA AAT ATC ATC ACC TGC ATT GAG      2064
Phe Val Pro Gly Lys Ser Ile Val Glu Asn Ile Ile Thr Cys Ile Glu
                655                 660                 665

AAG AGT TAC AAG TCC ATC TTT GTT TTG TCT CCC AAC TTT GTC CAG AGT      2112
Lys Ser Tyr Lys Ser Ile Phe Val Leu Ser Pro Asn Phe Val Gln Ser
            670                 675                 680

GAA TGG TGC CAT TAT GAA CTC TAC TTT GCC CAT CAC AAT CTC TTT CAT      2160
Glu Trp Cys His Tyr Glu Leu Tyr Phe Ala His His Asn Leu Phe His
        685                 690                 695

GAA GGA TCT AAT AGC TTA ATC CTG ATC TTG CTG GAA CCC ATT CCG CAG      2208
Glu Gly Ser Asn Ser Leu Ile Leu Ile Leu Leu Glu Pro Ile Pro Gln
    700                 705                 710

TAC TCC ATT CCT AGC AGT TAT CAC AAG CTC AAA AGT CTC ATG GCC AGG      2256
Tyr Ser Ile Pro Ser Ser Tyr His Lys Leu Lys Ser Leu Met Ala Arg
715                 720                 725                 730

AGG ACT TAT TTG GAA TGG CCC AAG GAA AAG AGC AAA CGT GGC CTT TTT      2304
Arg Thr Tyr Leu Glu Trp Pro Lys Glu Lys Ser Lys Arg Gly Leu Phe
                735                 740                 745

TGG GCT AAC TTA AGG GCA GCC ATT AAT ATT AAG CTG ACA GAG CAA GCA      2352
Trp Ala Asn Leu Arg Ala Ala Ile Asn Ile Lys Leu Thr Glu Gln Ala
            750                 755                 760

AAG AAA TAGTCTAGA                                                   2367
Lys Lys
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MTSIFHFAIIPMLILQIRIQLSEESEFLVDRSKNGLIHVPKDLSQKTTILNISQNYISELWTSDILSLSKLRILI
ISHNRIQYLDISVFKFNQELEYLDLSHNKLVKISCHPTVNLKHLDLSFNAFDALPICKEFGNMSQLKFLGLSTTH
LEKSSVLPIAHLNISKVLLVLGETYGEKEDPEGLQDFNTESLHIVFPTNKEFHFILDVSVKTVANLELSNIKCVL
EDNKCSYFLSILAKLQTNPKLSSLTLNNIETTWNSFIRILQLVWHTTVWYFSISNVKLQGQLDFRDFDYSGTSLK
ALSIHQVVSDVFGFPQSYIYEIFSNMNIKNFTVSGTRMVHMLCPSKISPFLHLDFSNNLLTDTVFENCGHLTELE
TLILQMNQLKELSKIAEMTTQMKSLQQLDISQNSVSYDEKKGDCSWTKSLLSLNMSSNILTDTIFRCLPPRIKVL
DLESNKIKSIPKQVVKLEALQELNVAFNSLTDLPGCGSFSSLSVLIIDHNSVSHPSADFFQSCQKMRSIKAGDNP
FQCTCELGEFVKNIDQVSSEVLEGWPDSYKCDYPESYRGTLLKDFHMSELSCNITLLIVTIVATMLVLAVTVTSL
CIYLDLPWYLRMVCQWTQTRRRARNIPLEELQRNLQFHAFISYSGHDSFWVKNELLPNLEKEGMQICLHERNFVP
GKSIVENIITCIEKSYKSIFVLSPNFVQSEWCHYELYFAHHNLFHEGSNSLILILLEPIPQYSIPSSYHKLKSLM
ARRTYLEWPKEKSKRGLFWANLRAAINIKLTEQAKK

Table 2: Nucleotide and amino acid sequences (see SEQ ID NO: 3 and 4) of a primate, e.g., human, DNAX Toll like Receptor 2 (DTLR2).

```
ATG CCA CAT ACT TTG TGG ATG GTG TGG GTC TTG GGG GTC ATC ATC AGC        48
Met Pro His Thr Leu Trp Met Val Trp Val Leu Gly Val Ile Ile Ser
-22     -20                 -15                 -10

CTC TCC AAG GAA GAA TCC TCC AAT CAG GCT TCT CTG TCT TGT GAC CGC        96
Leu Ser Lys Glu Glu Ser Ser Asn Gln Ala Ser Leu Ser Cys Asp Arg
     -5                  1                   5                  10

AAT GGT ATC TGC AAG GGC AGC TCA GGA TCT TTA AAC TCC ATT CCC TCA       144
Asn Gly Ile Cys Lys Gly Ser Ser Gly Ser Leu Asn Ser Ile Pro Ser
                 15                  20                  25

GGG CTC ACA GAA GCT GTA AAA AGC CTT GAC CTG TCC AAC AAC AGG ATC       192
Gly Leu Thr Glu Ala Val Lys Ser Leu Asp Leu Ser Asn Asn Arg Ile
             30                  35                  40

ACC TAC ATT AGC AAC AGT GAC CTA CAG AGG TGT GTG AAC CTC CAG GCT       240
Thr Tyr Ile Ser Asn Ser Asp Leu Gln Arg Cys Val Asn Leu Gln Ala
         45                  50                  55

CTG GTG CTG ACA TCC AAT GGA ATT AAC ACA ATA GAG GAA GAT TCT TTT       288
Leu Val Leu Thr Ser Asn Gly Ile Asn Thr Ile Glu Glu Asp Ser Phe
     60                  65                  70

TCT TCC CTG GGC AGT CTT GAA CAT TTA GAC TTA TCC TAT AAT TAC TTA       336
Ser Ser Leu Gly Ser Leu Glu His Leu Asp Leu Ser Tyr Asn Tyr Leu
 75                  80                  85                  90

TCT AAT TTA TCG TCT TCC TGG TTC AAG CCC CTT TCT TCT TTA ACA TTC       384
Ser Asn Leu Ser Ser Ser Trp Phe Lys Pro Leu Ser Ser Leu Thr Phe
                 95                 100                 105

TTA AAC TTA CTG GGA AAT CCT TAC AAA ACC CTA GGG GAA ACA TCT CTT       432
Leu Asn Leu Leu Gly Asn Pro Tyr Lys Thr Leu Gly Glu Thr Ser Leu
            110                 115                 120

TTT TCT CAT CTC ACA AAA TTG CAA ATC CTG AGA GTG GGA AAT ATG GAC       480
Phe Ser His Leu Thr Lys Leu Gln Ile Leu Arg Val Gly Asn Met Asp
        125                 130                 135

ACC TTC ACT AAG ATT CAA AGA AAA GAT TTT GCT GGA CTT ACC TTC CTT       528
Thr Phe Thr Lys Ile Gln Arg Lys Asp Phe Ala Gly Leu Thr Phe Leu
    140                 145                 150

GAG GAA CTT GAG ATT GAT GCT TCA GAT CTA CAG AGC TAT GAG CCA AAA       576
Glu Glu Leu Glu Ile Asp Ala Ser Asp Leu Gln Ser Tyr Glu Pro Lys
155                 160                 165                 170

AGT TTG AAG TCA ATT CAG AAC GTA AGT CAT CTG ATC CTT CAT ATG AAG       624
Ser Leu Lys Ser Ile Gln Asn Val Ser His Leu Ile Leu His Met Lys
                175                 180                 185
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CAG CAT ATT TTA CTG CTG GAG ATT TTT GTA GAT GTT ACA AGT TCC GTG      672
Gln His Ile Leu Leu Leu Glu Ile Phe Val Asp Val Thr Ser Ser Val
            190                 195                 200

GAA TGT TTG GAA CTG CGA GAT ACT GAT TTG GAC ACT TTC CAT TTT TCA      720
Glu Cys Leu Glu Leu Arg Asp Thr Asp Leu Asp Thr Phe His Phe Ser
        205                 210                 215

GAA CTA TCC ACT GGT GAA ACA AAT TCA TTG ATT AAA AAG TTT ACA TTT      768
Glu Leu Ser Thr Gly Glu Thr Asn Ser Leu Ile Lys Lys Phe Thr Phe
    220                 225                 230

AGA AAT GTG AAA ATC ACC GAT GAA AGT TTG TTT CAG GTT ATG AAA CTT      816
Arg Asn Val Lys Ile Thr Asp Glu Ser Leu Phe Gln Val Met Lys Leu
235                 240                 245                 250

TTG AAT CAG ATT TCT GGA CTG TTA GAA TTA GAG TTT GAT GAC TGT ACC      864
Leu Asn Gln Ile Ser Gly Leu Leu Glu Leu Glu Phe Asp Asp Cys Thr
                255                 260                 265

CTT AAT GGA GTT GGT AAT TTT AGA GCA TCT GAT AAT GAC AGA GTT ATA      912
Leu Asn Gly Val Gly Asn Phe Arg Ala Ser Asp Asn Asp Arg Val Ile
            270                 275                 280

GAT CCA GGT AAA GTG GAA ACG TTA ACA ATC CGG AGG CTG CAT ATT CCA      960
Asp Pro Gly Lys Val Glu Thr Leu Thr Ile Arg Arg Leu His Ile Pro
        285                 290                 295

AGG TTT TAC TTA TTT TAT GAT CTG AGC ACT TTA TAT TCA CTT ACA GAA     1008
Arg Phe Tyr Leu Phe Tyr Asp Leu Ser Thr Leu Tyr Ser Leu Thr Glu
    300                 305                 310

AGA GTT AAA AGA ATC ACA GTA GAA AAC AGT AAA GTT TTT CTG GTT CCT     1056
Arg Val Lys Arg Ile Thr Val Glu Asn Ser Lys Val Phe Leu Val Pro
315                 320                 325                 330

TGT TTA CTT TCA CAA CAT TTA AAA TCA TTA GAA TAC TTG GAT CTC AGT     1104
Cys Leu Leu Ser Gln His Leu Lys Ser Leu Glu Tyr Leu Asp Leu Ser
                335                 340                 345

GAA AAT TTG ATG GTT GAA GAA TAC TTG AAA AAT TCA GCC TGT GAG GAT     1152
Glu Asn Leu Met Val Glu Glu Tyr Leu Lys Asn Ser Ala Cys Glu Asp
            350                 355                 360

GCC TGG CCC TCT CTA CAA ACT TTA ATT TTA AGG CAA AAT CAT TTG GCA     1200
Ala Trp Pro Ser Leu Gln Thr Leu Ile Leu Arg Gln Asn His Leu Ala
        365                 370                 375

TCA TTG GAA AAA ACC GGA GAG ACT TTG CTC ACT CTG AAA AAC TTG ACT     1248
Ser Leu Glu Lys Thr Gly Glu Thr Leu Leu Thr Leu Lys Asn Leu Thr
    380                 385                 390

AAC ATT GAT ATC AGT AAG AAT AGT TTT CAT TCT ATG CCT GAA ACT TGT     1296
Asn Ile Asp Ile Ser Lys Asn Ser Phe His Ser Met Pro Glu Thr Cys
395                 400                 405                 410
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CAG TGG CCA GAA AAG ATG AAA TAT TTG AAC TTA TCC AGC ACA CGA ATA      1344
Gln Trp Pro Glu Lys Met Lys Tyr Leu Asn Leu Ser Ser Thr Arg Ile
                415                 420                 425

CAC AGT GTA ACA GGC TGC ATT CCC AAG ACA CTG GAA ATT TTA GAT GTT      1392
His Ser Val Thr Gly Cys Ile Pro Lys Thr Leu Glu Ile Leu Asp Val
            430                 435                 440

AGC AAC AAC AAT CTC AAT TTA TTT TCT TTG AAT TTG CCG CAA CTC AAA      1440
Ser Asn Asn Asn Leu Asn Leu Phe Ser Leu Asn Leu Pro Gln Leu Lys
        445                 450                 455

GAA CTT TAT ATT TCC AGA AAT AAG TTG ATG ACT CTA CCA GAT GCC TCC      1488
Glu Leu Tyr Ile Ser Arg Asn Lys Leu Met Thr Leu Pro Asp Ala Ser
    460                 465                 470

CTC TTA CCC ATG TTA CTA GTA TTG AAA ATC AGT AGG AAT GCA ATA ACT      1536
Leu Leu Pro Met Leu Leu Val Leu Lys Ile Ser Arg Asn Ala Ile Thr
475                 480                 485                 490

ACG TTT TCT AAG GAG CAA CTT GAC TCA TTT CAC ACA CTG AAG ACT TTG      1584
Thr Phe Ser Lys Glu Gln Leu Asp Ser Phe His Thr Leu Lys Thr Leu
                495                 500                 505

GAA GCT GGT GGC AAT AAC TTC ATT TGC TCC TGT GAA TTC CTC TCC TTC      1632
Glu Ala Gly Gly Asn Asn Phe Ile Cys Ser Cys Glu Phe Leu Ser Phe
            510                 515                 520

ACT CAG GAG CAG CAA GCA CTG GCC AAA GTC TTG ATT GAT TGG CCA GCA      1680
Thr Gln Glu Gln Gln Ala Leu Ala Lys Val Leu Ile Asp Trp Pro Ala
        525                 530                 535

AAT TAC CTG TGT GAC TCT CCA TCC CAT GTG CGT GGC CAG CAG GTT CAG      1728
Asn Tyr Leu Cys Asp Ser Pro Ser His Val Arg Gly Gln Gln Val Gln
    540                 545                 550

GAT GTC CGC CTC TCG GTG TCG GAA TGT CAC AGG ACA GCA CTG GTG TCT      1776
Asp Val Arg Leu Ser Val Ser Glu Cys His Arg Thr Ala Leu Val Ser
555                 560                 565                 570

GGC ATG TGC TGT GCT CTG TTC CTG CTG ATC CTG CTC ACG GGG GTC CTG      1824
Gly Met Cys Cys Ala Leu Phe Leu Leu Ile Leu Leu Thr Gly Val Leu
                575                 580                 585

TGC CAC CGT TTC CAT GGC CTG TGG TAT ATG AAA ATG ATG TGG GCC TGG      1872
Cys His Arg Phe His Gly Leu Trp Tyr Met Lys Met Met Trp Ala Trp
            590                 595                 600

CTC CAG GCC AAA AGG AAG CCC AGG AAA GCT CCC AGC AGG AAC ATC TGC      1920
Leu Gln Ala Lys Arg Lys Pro Arg Lys Ala Pro Ser Arg Asn Ile Cys
        605                 610                 615
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TAT GAT GCA TTT GTT TCT TAC AGT GAG CGG GAT GCC TAC TGG GTG GAG      1968
Tyr Asp Ala Phe Val Ser Tyr Ser Glu Arg Asp Ala Tyr Trp Val Glu
    620                 625                 630

AAC CTT ATG GTC CAG GAG CTG GAG AAC TTC AAT CCC CCC TTC AAG TTG      2016
Asn Leu Met Val Gln Glu Leu Glu Asn Phe Asn Pro Pro Phe Lys Leu
635                 640                 645                 650

TGT CTT CAT AAG CGG GAC TTC ATT CCT GGC AAG TGG ATC ATT GAC AAT      2064
Cys Leu His Lys Arg Asp Phe Ile Pro Gly Lys Trp Ile Ile Asp Asn
                655                 660                 665

ATC ATT GAC TCC ATT GAA AAG AGC CAC AAA ACT GTC TTT GTG CTT TCT      2112
Ile Ile Asp Ser Ile Glu Lys Ser His Lys Thr Val Phe Val Leu Ser
            670                 675                 680

GAA AAC TTT GTG AAG AGT GAG TGG TGC AAG TAT GAA CTG GAC TTC TCC      2160
Glu Asn Phe Val Lys Ser Glu Trp Cys Lys Tyr Glu Leu Asp Phe Ser
        685                 690                 695

CAT TTC CGT CTT TTT GAA GAG AAC AAT GAT GCT GCC ATT CTC ATT CTT      2208
His Phe Arg Leu Phe Glu Glu Asn Asn Asp Ala Ala Ile Leu Ile Leu
    700                 705                 710

CTG GAG CCC ATT GAG AAA AAA GCC ATT CCC CAG CGC TTC TGC AAG CTG      2256
Leu Glu Pro Ile Glu Lys Lys Ala Ile Pro Gln Arg Phe Cys Lys Leu
715                 720                 725                 730

CGG AAG ATA ATG AAC ACC AAG ACC TAC CTG GAG TGG CCC ATG GAC GAG      2304
Arg Lys Ile Met Asn Thr Lys Thr Tyr Leu Glu Trp Pro Met Asp Glu
                735                 740                 745

GCT CAG CGC GAA GGA TTT TGG GTA AAT CTG AGA GCT GCG ATA AAG TCC      2352
Ala Gln Arg Glu Gly Phe Trp Val Asn Leu Arg Ala Ala Ile Lys Ser
            750                 755                 760

TAG                                                                  2355
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MPHTLWMVWVLGVIISLSKEESSNQASLSCDRNGICKGSSGSLNSIPSGLTEAVKSLDLSNNRITYISNSDLQRC
VNLQALVLTSNGINTIEEDSFSSLGSLEHLDLSYNYLSNLSSSWFKPLSSLTFLNLLGNPYKTLGETSLFSHLTK
LQILRVGNMDTFTKIQRKDFAGLTFLEELEIDASDLQSYEPKSLKSIQNVSHLILHMKQHILLLEIFVDVTSSVE
CLELRDTDLDTFHFSELSTGETNSLIKKFTFRNVKITDESLFQVMKLLNQISGLLELEFDDCTLNGVGNFRASDN
DRVIDPGKVETLTIRRLHIPRFYLFYDLSTLYSLTERVKRITVENSKVFLVPCLLSQHLKSLEYLDLSENLMVEE
YLKNSACEDAWPSLQTLILRQNHLASLEKTGETLLTLKNLTNIDISKNSFHSMPETCQWPEKMKYLNLSSTRIHS
VTGCIPKTLEILDVSNNNLNLFSLNLPQLKELYISRNKLMTLPDASLLPMLLVLKISRNAITTFSKEQLDSFHTL
KTLEAGGNNFICSCEFLSFIQEQQALAKVLIDWPANYLCDSPSHVRGQQVQDVRLSVSECHRTALVSGMCCALFL
LILLTGVLCHRFHGLWYMKMMWAWLQAKRKPRKAPSRNICYDAFVSYSERDAYWVENLMVQELENFNPPFKLCLH
KRDFIPGKWIIDNIIDSIEKSHKTVFVLSENFVKSEWCKYELDFSHFRLFEENNDAAILILLEPIEKKAIPQRFC
KLRKIMNIKTYLEWPMDEAQREGFWVNLRAAIKS
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Table 3: Nucleotide and amino acid sequences (see SEQ ID NO: 5 and 6) of a mammalian, e.g., human, Toll like Receptor 3 (DTLR3).

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ATG AGA CAG ACT TTG CCT TGT ATC TAC TTT TGG GGG GGC CTT TTG CCC        48
Met Arg Gln Thr Leu Pro Cys Ile Tyr Phe Trp Gly Gly Leu Leu Pro
-21 -20                 -15                 -10

TTT GGG ATG CTG TGT GCA TCC TCC ACC ACC AAG TGC ACT GTT AGC CAT        96
Phe Gly Met Leu Cys Ala Ser Ser Thr Thr Lys Cys Thr Val Ser His
 -5                   1                   5                  10

GAA GTT GCT GAC TGC AGC CAC CTG AAG TTG ACT CAG GTA CCC GAT GAT       144
Glu Val Ala Asp Cys Ser His Leu Lys Leu Thr Gln Val Pro Asp Asp
               15                  20                  25

CTA CCC ACA AAC ATA ACA GTG TTG AAC CTT ACC CAT AAT CAA CTC AGA       192
Leu Pro Thr Asn Ile Thr Val Leu Asn Leu Thr His Asn Gln Leu Arg
          30                  35                  40

AGA TTA CCA GCC GCC AAC TTC ACA AGG TAT AGC CAG CTA ACT AGC TTG       240
Arg Leu Pro Ala Ala Asn Phe Thr Arg Tyr Ser Gln Leu Thr Ser Leu
     45                  50                  55

GAT GTA GGA TTT AAC ACC ATC TCA AAA CTG GAG CCA GAA TTG TGC CAG       288
Asp Val Gly Phe Asn Thr Ile Ser Lys Leu Glu Pro Glu Leu Cys Gln
 60                  65                  70                  75

AAA CTT CCC ATG TTA AAA GTT TTG AAC CTC CAG CAC AAT GAG CTA TCT       336
Lys Leu Pro Met Leu Lys Val Leu Asn Leu Gln His Asn Glu Leu Ser
                 80                  85                  90

CAA CTT TCT GAT AAA ACC TTT GCC TTC TGC ACG AAT TTG ACT GAA CTC       384
Gln Leu Ser Asp Lys Thr Phe Ala Phe Cys Thr Asn Leu Thr Glu Leu
               95                 100                 105

CAT CTC ATG TCC AAC TCA ATC CAG AAA ATT AAA AAT AAT CCC TTT GTC       432
His Leu Met Ser Asn Ser Ile Gln Lys Ile Lys Asn Asn Pro Phe Val
          110                 115                 120

AAG CAG AAG AAT TTA ATC ACA TTA GAT CTG TCT CAT AAT GGC TTG TCA       480
Lys Gln Lys Asn Leu Ile Thr Leu Asp Leu Ser His Asn Gly Leu Ser
     125                 130                 135

TCT ACA AAA TTA GGA ACT CAG GTT CAG CTG GAA AAT CTC CAA GAG CTT       528
Ser Thr Lys Leu Gly Thr Gln Val Gln Leu Glu Asn Leu Gln Glu Leu
140                 145                 150                 155

CTA TTA TCA AAC AAT AAA ATT CAA GCG CTA AAA AGT GAA GAA CTG GAT       576
Leu Leu Ser Asn Asn Lys Ile Gln Ala Leu Lys Ser Glu Glu Leu Asp
                160                 165                 170

ATC TTT GCC AAT TCA TCT TTA AAA AAA TTA GAG TTG TCA TCG AAT CAA       624
Ile Phe Ala Asn Ser Ser Leu Lys Lys Leu Glu Leu Ser Ser Asn Gln
               175                 180                 185
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ATT AAA GAG TTT TCT CCA GGG TGT TTT CAC GCA ATT GGA AGA TTA TTT      672
Ile Lys Glu Phe Ser Pro Gly Cys Phe His Ala Ile Gly Arg Leu Phe
        190                 195                 200

GGC CTC TTT CTG AAC AAT GTC CAG CTG GGT CCC AGC CTT ACA GAG AAG      720
Gly Leu Phe Leu Asn Asn Val Gln Leu Gly Pro Ser Leu Thr Glu Lys
    205                 210                 215

CTA TGT TTG GAA TTA GCA AAC ACA AGC ATT CGG AAT CTG TCT CTG AGT      768
Leu Cys Leu Glu Leu Ala Asn Thr Ser Ile Arg Asn Leu Ser Leu Ser
220                 225                 230                 235

AAC AGC CAG CTG TCC ACC ACC AGC AAT ACA ACT TTC TTG GGA CTA AAG      816
Asn Ser Gln Leu Ser Thr Thr Ser Asn Thr Thr Phe Leu Gly Leu Lys
                240                 245                 250

TGG ACA AAT CTC ACT ATG CTC GAT CTT TCC TAC AAC AAC TTA AAT GTG      864
Trp Thr Asn Leu Thr Met Leu Asp Leu Ser Tyr Asn Asn Leu Asn Val
            255                 260                 265

GTT GGT AAC GAT TCC TTT GCT TGG CTT CCA CAA CTA GAA TAT TTC TTC      912
Val Gly Asn Asp Ser Phe Ala Trp Leu Pro Gln Leu Glu Tyr Phe Phe
        270                 275                 280

CTA GAG TAT AAT AAT ATA CAG CAT TTG TTT TCT CAC TCT TTG CAC GGG      960
Leu Glu Tyr Asn Asn Ile Gln His Leu Phe Ser His Ser Leu His Gly
    285                 290                 295

CTT TTC AAT GTG AGG TAC CTG AAT TTG AAA CGG TCT TCT ACT AAA CAA     1008
Leu Phe Asn Val Arg Tyr Leu Asn Leu Lys Arg Ser Phe Thr Lys Gln
300                 305                 310                 315

AGT ATT TCC CTT GCC TCA CTC CCC AAG ATT GAT GAT TTT TCT TTT CAG     1056
Ser Ile Ser Leu Ala Ser Leu Pro Lys Ile Asp Asp Phe Ser Phe Gln
                320                 325                 330

TGG CTA AAA TGT TTG GAG CAC CTT AAC ATG GAA GAT AAT GAT ATT CCA     1104
Trp Leu Lys Cys Leu Glu His Leu Asn Met Glu Asp Asn Asp Ile Pro
            335                 340                 345

GGC ATA AAA AGC AAT ATG TTC ACA GGA TTG ATA AAC CTG AAA TAC TTA     1152
Gly Ile Lys Ser Asn Met Phe Thr Gly Leu Ile Asn Leu Lys Tyr Leu
        350                 355                 360

AGT CTA TCC AAC TCC TTT ACA AGT TTG CGA ACT TTG ACA AAT GAA ACA     1200
Ser Leu Ser Asn Ser Phe Thr Ser Leu Arg Thr Leu Thr Asn Glu Thr
    365                 370                 375

TTT GTA TCA CTT GCT CAT TCT CCC TTA CAC ATA CTC AAC CTA ACC AAG     1248
Phe Val Ser Leu Ala His Ser Pro Leu His Ile Leu Asn Leu Thr Lys
380                 385                 390                 395

AAT AAA ATC TCA AAA ATA GAG AGT GAT GCT TTC TCT TGG TTG GGC CAC     1296
Asn Lys Ile Ser Lys Ile Glu Ser Asp Ala Phe Ser Trp Leu Gly His
                400                 405                 410
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CTA GAA GTA CTT GAC CTG GGC CTT AAT GAA ATT GGG CAA GAA CTC ACA   1344
Leu Glu Val Leu Asp Leu Gly Leu Asn Glu Ile Gly Gln Glu Leu Thr
            415                 420                 425

GGC CAG GAA TGG AGA GGT CTA GAA AAT ATT TTC GAA ATC TAT CTT TCC   1392
Gly Gln Glu Trp Arg Gly Leu Glu Asn Ile Phe Glu Ile Tyr Leu Ser
        430                 435                 440

TAC AAC AAG TAC CTG CAG CTG ACT AGG AAC TCC TTT GCC TTG GTC CCA   1440
Tyr Asn Lys Tyr Leu Gln Leu Thr Arg Asn Ser Phe Ala Leu Val Pro
    445                 450                 455

AGC CTT CAA CGA CTG ATG CTC CGA ACG GTG GCC CTT AAA AAT GTG GAT   1488
Ser Leu Gln Arg Leu Met Leu Arg Arg Val Ala Leu Lys Asn Val Asp
460                 465                 470                 475

AGC TCT CCT TCA CCA TTC CAG CCT CTT CGT AAC TTG ACC ATC CTG GAT   1536
Ser Ser Pro Ser Pro Phe Gln Pro Leu Arg Asn Leu Thr Ile Leu Asp
                480                 485                 490

CTA AGC AAC AAC AAC ATA GCC AAC ATA AAT GAT GAC ATG TTG GAG GGT   1584
Leu Ser Asn Asn Asn Ile Ala Asn Ile Asn Asp Asp Met Leu Glu Gly
            495                 500                 505

CTT GAG AAA CTA GAA ATT CTC GAT TTG CAG CAT AAC AAC TTA GCA CGG   1632
Leu Glu Lys Leu Glu Ile Leu Asp Leu Gln His Asn Asn Leu Ala Arg
        510                 515                 520

CTC TGG AAA CAC GCA AAC CCT GGT GGT CCC ATT TAT TTC CTA AAG GGT   1680
Leu Trp Lys His Ala Asn Pro Gly Gly Pro Ile Tyr Phe Leu Lys Gly
    525                 530                 535

CTG TCT CAC CTC CAC ATC CTT AAC TTG GAG TCC AAC GGC TTT GAC GAG   1728
Leu Ser His Leu His Ile Leu Asn Leu Glu Ser Asn Gly Phe Asp Glu
540                 545                 550                 555

ATC CCA GTT GAG GTC TTC AAG GAT TTA TTT GAA CTA AAG ATC ATC GAT   1776
Ile Pro Val Glu Val Phe Lys Asp Leu Phe Glu Leu Lys Ile Ile Asp
                560                 565                 570

TTA GGA TTG AAT AAT TTA AAC ACA CTT CCA GCA TCT GTC TTT AAT AAT   1824
Leu Gly Leu Asn Asn Leu Asn Thr Leu Pro Ala Ser Val Phe Asn Asn
            575                 580                 585

CAG GTG TCT CTA AAG TCA TTG AAC CTT CAG AAG AAT CTC ATA ACA TCC   1872
Gln Val Ser Leu Lys Ser Leu Asn Leu Gln Lys Asn Leu Ile Thr Ser
        590                 595                 600

GTT GAG AAG AAG GTT TTC GGG CCA GCT TTC AGG AAC CTG ACT GAG TTA   1920
Val Glu Lys Lys Val Phe Gly Pro Ala Phe Arg Asn Leu Thr Glu Leu
    605                 610                 615
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GAT ATG CGC TTT AAT CCC TTT GAT TGC ACG TGT GAA AGT ATT GCC TGG      1968
Asp Met Arg Phe Asn Pro Phe Asp Cys Thr Cys Glu Ser Ile Ala Trp
620                 625                 630                 635

TTT GTT AAT TGG ATT AAC GAG ACC CAT ACC AAC ATC CCT GAG CTG TCA      2016
Phe Val Asn Trp Ile Asn Glu Thr His Thr Asn Ile Pro Glu Leu Ser
                640                 645                 650

AGC CAC TAC CTT TGC AAC ACT CCA CCT CAC TAT CAT GGG TTC CCA GTG      2064
Ser His Tyr Leu Cys Asn Thr Pro Pro His Tyr His Gly Phe Pro Val
            655                 660                 665

AGA CTT TTT GAT ACA TCA TCT TGC AAA GAC AGT GCC CCC TTT GAA CTC      2112
Arg Leu Phe Asp Thr Ser Ser Cys Lys Asp Ser Ala Pro Phe Glu Leu
        670                 675                 680

TTT TTC ATG ATC AAT ACC AGT ATC CTG TTG ATT TTT ATC TTT ATT GTA      2160
Phe Phe Met Ile Asn Thr Ser Ile Leu Leu Ile Phe Ile Phe Ile Val
    685                 690                 695

CTT CTC ATC CAC TTT GAG GGC TGG AGG ATA TCT TTT TAT TGG AAT GTT      2208
Leu Leu Ile His Phe Glu Gly Trp Arg Ile Ser Phe Tyr Trp Asn Val
700                 705                 710                 715

TCA GTA CAT CGA GTT CTT GGT TTC AAA GAA ATA GAC AGA CAG ACA GAA      2256
Ser Val His Arg Val Leu Gly Phe Lys Glu Ile Asp Arg Gln Thr Glu
                720                 725                 730

CAG TTT GAA TAT GCA GCA TAT ATA ATT CAT GCC TAT AAA GAT AAG GAT      2304
Gln Phe Glu Tyr Ala Ala Tyr Ile Ile His Ala Tyr Lys Asp Lys Asp
            735                 740                 745

TGG GTC TGG GAA CAT TTC TCT TCA ATG GAA AAG GAA GAC CAA TCT CTC      2352
Trp Val Trp Glu His Phe Ser Ser Met Glu Lys Glu Asp Gln Ser Leu
        750                 755                 760

AAA TTT TGT CTG GAA GAA AGG GAC TTT GAG GCG GGT GTT TTT GAA CTA      2400
Lys Phe Cys Leu Glu Glu Arg Asp Phe Glu Ala Gly Val Phe Glu Leu
    765                 770                 775

GAA GCA ATT GTT AAC AGC ATC AAA AGA AGC AGA AAA ATT ATT TTT GTT      2448
Glu Ala Ile Val Asn Ser Ile Lys Arg Ser Arg Lys Ile Ile Phe Val
780                 785                 790                 795

ATA ACA CAC CAT CTA TTA AAA GAC CCA TTA TGC AAA AGA TTC AAG GTA      2496
Ile Thr His His Leu Leu Lys Asp Pro Leu Cys Lys Arg Phe Lys Val
                800                 805                 810

CAT CAT GCA GTT CAA CAA GCT ATT GAA CAA AAT CTG GAT TCC ATT ATA      2544
His His Ala Val Gln Gln Ala Ile Glu Gln Asn Leu Asp Ser Ile Ile
            815                 820                 825

TTG GTT TTC CTT GAG GAG ATT CCA GAT TAT AAA CTG AAC CAT GCA CTC      2592
Leu Val Phe Leu Glu Glu Ile Pro Asp Tyr Lys Leu Asn His Ala Leu
        830                 835                 840
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TGT TTG CGA AGA GGA ATG TTT AAA TCT CAC TGC ATC TTG AAC TGG CCA      2640
Cys Leu Arg Arg Gly Met Phe Lys Ser His Cys Ile Leu Asn Trp Pro
    845                 850                 855

GTT CAG AAA GAA CGG ATA GGT GCC TTT CGT CAT AAA TTG CAA GTA GCA      2688
Val Gln Lys Glu Arg Ile Gly Ala Phe Arg His Lys Leu Gln Val Ala
860                 865                 870                 875

CTT GGA TCC AAA AAC TCT GTA CAT TAA                                  2715
Leu Gly Ser Lys Asn Ser Val His
                880
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MRQTLPCIYFWGGLLPFGMICASSTTKCTVSHEVADCSHLKLTQVPDDLPTNITVLNLTHNQLRRLPAANFTRYS
QLTSLDVGFNTISKLEPELCQKLPMLKVLNLQHNELSQLSDKTFAFCTNLTELHLMSNSIQKIKNNPFVKQKNLI
TLDLSHNGLSSTKLGTQVQLENLQELLLSNNKIQALKSEELDIFANSSLKKLELSSNQIKEFSPGCFHAIGRLFG
LFLNNVQLGPSLTEKLCLELANTSIRNLSLSNSQLSTTSNTTFLGLKWTNLTMLDLSYNNLNVVGNDSFAWLPQL
EYFFLEYNNIQHLFSHSLHGLFNVRYLNLKRSFTKQSISLASLPKIDDFSFQWLKCLEHLNMEDNDIPGIKSNMF
TGLINLKYLSLSNSFTSLRTLTNETFVSLAHSPLHILNLTKNKISKIESDAFSWLGHLEVLDLGLNEIGQELTGQ
EWRGLENIFEIYLSYNKYLQLTRNSFALVPSLQRLMLRRVALKNVDSSPSPFQPLRNLTILDLSNNNIANINDDM
LEGLEKLEILDLQHNNLARLWKHANPGGPIYFLKGLSHLHILNLESNGFDEIPVEVFKDLFELKIIDLGLNNLNT
LPASVFNNQVSLKSLNLQKNLITSVEKKVFGPAFRNLTELDMRFNPFDCTCESIAWFVNWINETHTNIPELSSHY
LCNTPPHYHGFPVRLFDTSSCKDSAPFELFFMINTSILLIFIFIVLLIHFEGWRISFYWNVSVHRVLGFKEIDRQ
TEQFEYAAYIIHAYKDKDWVWEHFSSMEKEDQSLKFCLEERDFEAGVFELEAIVNSIKRSRKIIFVITHHLLKDP
LCKRFKVHHAVQQAIEQNLDSIILVFLEEIPDYKLNHALCLRRGMFKSHCILNWPVQKERIGAFRHKLQVALGSK
NSVH

Table 4: Nucleotide and amino acid sequences (see SEQ ID NO: 7 and 8) of a mammalian, e.g., primate, human, DNAX Toll like Receptor 4 (DTLR4).

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ATG GAG CTG AAT TTC TAC AAA ATC CCC GAC AAC CTC CCC TTC TCA ACC      48
Met Glu Leu Asn Phe Tyr Lys Ile Pro Asp Asn Leu Pro Phe Ser Thr
 1               5                  10                  15

AAG AAC CTG GAC CTG AGC TTT AAT CCC CTG AGG CAT TTA GGC AGC TAT      96
Lys Asn Leu Asp Leu Ser Phe Asn Pro Leu Arg His Leu Gly Ser Tyr
            20                  25                  30

AGC TTC TTC AGT TTC CCA GAA CTG CAG GTG CTG GAT TTA TCC AGG TGT     144
Ser Phe Phe Ser Phe Pro Glu Leu Gln Val Leu Asp Leu Ser Arg Cys
        35                  40                  45

GAA ATC CAG ACA ATT GAA GAT GGG GCA TAT CAG AGC CTA AGC CAC CTC     192
Glu Ile Gln Thr Ile Glu Asp Gly Ala Tyr Gln Ser Leu Ser His Leu
    50                  55                  60

TCT ACC TTA ATA TTG ACA GGA AAC CCC ATC CAG AGT TTA GCC CTG GGA     240
Ser Thr Leu Ile Leu Thr Gly Asn Pro Ile Gln Ser Leu Ala Leu Gly
65                  70                  75                  80

GCC TTT TCT GGA CTA TCA AGT TTA CAG AAG CTG GTG GCT GTG GAG ACA     288
Ala Phe Ser Gly Leu Ser Ser Leu Gln Lys Leu Val Ala Val Glu Thr
                85                  90                  95

AAT CTA GCA TCT CTA GAG AAC TTC CCC ATT GGA CAT CTC AAA ACT TTG     336
Asn Leu Ala Ser Leu Glu Asn Phe Pro Ile Gly His Leu Lys Thr Leu
            100                 105                 110

AAA GAA CTT AAT GTG GCT CAC AAT CTT ATC CAA TCT TTC AAA TTA CCT     384
Lys Glu Leu Asn Val Ala His Asn Leu Ile Gln Ser Phe Lys Leu Pro
        115                 120                 125

GAG TAT TTT TCT AAT CTG ACC AAT CTA GAG CAC TTG GAC CTT TCC AGC     432
Glu Tyr Phe Ser Asn Leu Thr Asn Leu Glu His Leu Asp Leu Ser Ser
    130                 135                 140

AAC AAG ATT CAA AGT ATT TAT TGC ACA GAC TTG CGG GTT CTA CAT CAA     480
Asn Lys Ile Gln Ser Ile Tyr Cys Thr Asp Leu Arg Val Leu His Gln
145                 150                 155                 160

ATG CCC CTA CTC AAT CTC TCT TTA GAC CTG TCC CTG AAC CCT ATG AAC     528
Met Pro Leu Leu Asn Leu Ser Leu Asp Leu Ser Leu Asn Pro Met Asn
                165                 170                 175

TTT ATC CAA CCA GGT GCA TTT AAA GAA ATT AGG CTT CAT AAG CTG ACT     576
Phe Ile Gln Pro Gly Ala Phe Lys Glu Ile Arg Leu His Lys Leu Thr
            180                 185                 190

TTA AGA AAT AAT TTT GAT AGT TTA AAT GTA ATG AAA ACT TGT ATT CAA     624
Leu Arg Asn Asn Phe Asp Ser Leu Asn Val Met Lys Thr Cys Ile Gln
        195                 200                 205
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GGT CTG GCT GGT TTA GAA GTC CAT CGT TTG GTT CTG GGA GAA TTT AGA      672
Gly Leu Ala Gly Leu Glu Val His Arg Leu Val Leu Gly Glu Phe Arg
    210                 215                 220

AAT GAA GGA AAC TTG GAA AAG TTT GAC AAA TCT GCT CTA GAG GGC CTG      720
Asn Glu Gly Asn Leu Glu Lys Phe Asp Lys Ser Ala Leu Glu Gly Leu
225                 230                 235                 240

TGC AAT TTG ACC ATT GAA GAA TTC CGA TTA GCA TAC TTA GAC TAC TAC      768
Cys Asn Leu Thr Ile Glu Glu Phe Arg Leu Ala Tyr Leu Asp Tyr Tyr
                245                 250                 255

CTC GAT GAT ATT ATT GAC TTA TTT AAT TGT TTG ACA AAT GTT TCT TCA      816
Leu Asp Asp Ile Ile Asp Leu Phe Asn Cys Leu Thr Asn Val Ser Ser
            260                 265                 270

TTT TCC CTG GTG AGT GTG ACT ATT GAA AGG GTA AAA GAC TTT TCT TAT      864
Phe Ser Leu Val Ser Val Thr Ile Glu Arg Val Lys Asp Phe Ser Tyr
        275                 280                 285

AAT TTC GGA TGG CAA CAT TTA GAA TTA GTT AAC TGT AAA TTT GGA CAG      912
Asn Phe Gly Trp Gln His Leu Glu Leu Val Asn Cys Lys Phe Gly Gln
    290                 295                 300

TTT CCC ACA TTG AAA CTC AAA TCT CTC AAA AGG CTT ACT TTC ACT TCC      960
Phe Pro Thr Leu Lys Leu Lys Ser Leu Lys Arg Leu Thr Phe Thr Ser
305                 310                 315                 320

AAC AAA GGT GGG AAT GCT TTT TCA GAA GTT GAT CTA CCA AGC CTT GAG     1008
Asn Lys Gly Gly Asn Ala Phe Ser Glu Val Asp Leu Pro Ser Leu Glu
                325                 330                 335

TTT CTA GAT CTC AGT AGA AAT GGC TTG AGT TTC AAA GGT TGC TGT TCT     1056
Phe Leu Asp Leu Ser Arg Asn Gly Leu Ser Phe Lys Gly Cys Cys Ser
            340                 345                 350

CAA AGT GAT TTT GGG ACA ACC AGC CTA AAG TAT TTA GAT CTG AGC TTC     1104
Gln Ser Asp Phe Gly Thr Thr Ser Leu Lys Tyr Leu Asp Leu Ser Phe
        355                 360                 365

AAT GGT GTT ATT ACC ATG AGT TCA AAC TTC TTG GGC TTA GAA CAA CTA     1152
Asn Gly Val Ile Thr Met Ser Ser Asn Phe Leu Gly Leu Glu Gln Leu
    370                 375                 380

GAA CAT CTG GAT TTC CAG CAT TCC AAT TTG AAA CAA ATG AGT GAG TTT     1200
Glu His Leu Asp Phe Gln His Ser Asn Leu Lys Gln Met Ser Glu Phe
385                 390                 395                 400

TCA GTA TTC CTA TCA CTC AGA AAC CTC ATT TAC CTT GAC ATT TCT CAT     1248
Ser Val Phe Leu Ser Leu Arg Asn Leu Ile Tyr Leu Asp Ile Ser His
                405                 410                 415

ACT CAC ACC AGA GTT GCT TTC AAT GGC ATC TTC AAT GGC TTG TCC AGT     1296
Thr His Thr Arg Val Ala Phe Asn Gly Ile Phe Asn Gly Leu Ser Ser
            420                 425                 430
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CTC GAA GTC TTG AAA ATG GCT GGC AAT TCT TTC CAG GAA AAC TTC CTT      1344
Leu Glu Val Leu Lys Met Ala Gly Asn Ser Phe Gln Glu Asn Phe Leu
        435                 440                 445

CCA GAT ATC TTC ACA GAG CTG AGA AAC TTG ACC TTC CTG GAC CTC TCT      1392
Pro Asp Ile Phe Thr Glu Leu Arg Asn Leu Thr Phe Leu Asp Leu Ser
    450                 455                 460

CAG TGT CAA CTG GAG CAG TTG TCT CCA ACA GCA TTT AAC TCA CTC TCC      1440
Gln Cys Gln Leu Glu Gln Leu Ser Pro Thr Ala Phe Asn Ser Leu Ser
465                 470                 475                 480

AGT CTT CAG GTA CTA AAT ATG AGC CAC AAC AAC TTC TTT TCA TTG GAT      1488
Ser Leu Gln Val Leu Asn Met Ser His Asn Asn Phe Phe Ser Leu Asp
                485                 490                 495

ACG TTT CCT TAT AAG TGT CTG AAC TCC CTC CAG GTT CTT GAT TAC AGT      1536
Thr Phe Pro Tyr Lys Cys Leu Asn Ser Leu Gln Val Leu Asp Tyr Ser
            500                 505                 510

CTC AAT CAC ATA ATG ACT TCC AAA AAA CAG GAA CTA CAG CAT TTT CCA      1584
Leu Asn His Ile Met Thr Ser Lys Lys Gln Glu Leu Gln His Phe Pro
        515                 520                 525

AGT AGT CTA GCT TTC TTA AAT CTT ACT CAG AAT GAC TTT GCT TGT ACT      1632
Ser Ser Leu Ala Phe Leu Asn Leu Thr Gln Asn Asp Phe Ala Cys Thr
    530                 535                 540

TGT GAA CAC CAG AGT TTC CTG CAA TGG ATC AAG GAC CAG AGG CAG CTC      1680
Cys Glu His Gln Ser Phe Leu Gln Trp Ile Lys Asp Gln Arg Gln Leu
545                 550                 555                 560

TTG GTG GAA GTT GAA CGA ATG GAA TGT GCA ACA CCT TCA GAT AAG CAG      1728
Leu Val Glu Val Glu Arg Met Glu Cys Ala Thr Pro Ser Asp Lys Gln
                565                 570                 575

GGC ATG CCT GTG CTG AGT TTG AAT ATC ACC TGT CAG ATG AAT AAG ACC      1776
Gly Met Pro Val Leu Ser Leu Asn Ile Thr Cys Gln Met Asn Lys Thr
            580                 585                 590

ATC ATT GGT GTG TCG GTC CTC AGT GTG CTT GTA GTA TCT GTT GTA GCA      1824
Ile Ile Gly Val Ser Val Leu Ser Val Leu Val Val Ser Val Val Ala
        595                 600                 605

GTT CTG GTC TAT AAG TTC TAT TTT CAC CTG ATG CTT CTT GCT GGC TGC      1872
Val Leu Val Tyr Lys Phe Tyr Phe His Leu Met Leu Leu Ala Gly Cys
    610                 615                 620

ATA AAG TAT GGT AGA GGT GAA AAC ATC TAT GAT GCC TTT GTT ATC TAC      1920
Ile Lys Tyr Gly Arg Gly Glu Asn Ile Tyr Asp Ala Phe Val Ile Tyr
625                 630                 635                 640
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TCA AGC CAG GAT GAG GAC TGG GTA AGG AAT GAG CTA GTA AAG AAT TTA      1968
Ser Ser Gln Asp Glu Asp Trp Val Arg Asn Glu Leu Val Lys Asn Leu
                645                 650                 655

GAA GAA GGG GTG CCT CCA TTT CAG CTC TGC CTT CAC TAC AGA GAC TTT      2016
Glu Glu Gly Val Pro Pro Phe Gln Leu Cys Leu His Tyr Arg Asp Phe
            660                 665                 670

ATT CCC GGT GTG GCC ATT GCT GCC AAC ATC ATC CAT GAA GGT TTC CAT      2064
Ile Pro Gly Val Ala Ile Ala Ala Asn Ile Ile His Glu Gly Phe His
        675                 680                 685

AAA AGC CGA AAG GTG ATT GTT GTG GTG TCC CAG CAC TTC ATC CAG AGC      2112
Lys Ser Arg Lys Val Ile Val Val Val Ser Gln His Phe Ile Gln Ser
    690                 695                 700

CGC TGG TGT ATC TTT GAA TAT GAG ATT GCT CAG ACC TGG CAG TTT CTG      2160
Arg Trp Cys Ile Phe Glu Tyr Glu Ile Ala Gln Thr Trp Gln Phe Leu
705                 710                 715                 720

AGC AGC CGT GCT GGT ATC ATC TTC ATT GTC CTG CAG AAG GTG GAG AAG      2208
Ser Ser Arg Ala Gly Ile Ile Phe Ile Val Leu Gln Lys Val Glu Lys
                725                 730                 735

ACC CTG CTC AGG CAG CAG GTG GAG CTG TAC CGC CTT CTC AGC AGG AAC      2256
Thr Leu Leu Arg Gln Gln Val Glu Leu Tyr Arg Leu Leu Ser Arg Asn
            740                 745                 750

ACT TAC CTG GAG TGG GAG GAC AGT GTC CTG GGG CGG CAC ATC TTC TGG      2304
Thr Tyr Leu Glu Trp Glu Asp Ser Val Leu Gly Arg His Ile Phe Trp
        755                 760                 765

AGA CGA CTC AGA AAA GCC CTG CTG GAT GGT AAA TCA TGG AAT CCA GAA      2352
Arg Arg Leu Arg Lys Ala Leu Leu Asp Gly Lys Ser Trp Asn Pro Glu
    770                 775                 780

GGA ACA GTG GGT ACA GGA TGC AAT TGG CAG GAA GCA ACA TCT ATC          2397
Gly Thr Val Gly Thr Gly Cys Asn Trp Gln Glu Ala Thr Ser Ile
785                 790                 795

TGA                                                                 2400
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MELNFYKIPDNLPFSTKNLDLSFNPLRHLGSYSFFSFPELQVLDLSRCEIQTIEDGAYQSLSHLSTLILTGNP
IQSLALGAFSGLSSLQKLVAVETNLASLENFPIGHLKTLKELNVAHNLIQSFKLPEYFSNLTNLEHLDLSSNK
IQSIYCTDLRVLHQMPLLNLSLDLSLNPMNFIQPGAFKEIRLHKLTLRNNFDSLNVMKTCIQGLAGLEVHRLV
LGEFRNEGNLEKFDKSALEGLCNLTIEEFRLAYLDYYLDDIIDLFNCLTNVSSFSLVSVTIERVKDFSYNFGW
QHLELVNCKFGQFPTLKLKSLKRLTFTSNKGGNAFSEVDLPSLEFLDLSRNGLSFKGCCSQSDFGTTSLKYLD
LSFNGVITMSSNFLGLEQLEHLDFQHSNLKQMSEFSVFLSLRNLIYLDISHTHTRVAFNGIFNGLSSLEVLKM
AGNSFQENFLPDIFTELRNLTFLDLSQCQLEQLSPTAFNSLSSLQVLNMSHNNFFSLDTFPYKCLNSLQVLDY
SLNHIMTSKKQELQHFPSSLAFLNLTQNDFACTCEHQSFLQWIKDQRQLLVEVERMECATPSDKQGMPVLSLN
ITCQMNKTIIGVSVLSVLVVSVVAVLVYKFYFHLMLLAGCIKYGRGENIYDAFVIYSSQDEDWVRNELVKNLE
EGVPPFQLCLHYRDFIPGVAIAANIIHEGFHKSRKVIVVVSQHFIQSRWCIFEYEIAQTWQFLSSRAGIIFIV
LQKVEKTLLRQQVELYRLLSRNTYLEWEDSVLGRHIFWRRLRKALLDGKSWNPEGTVGTGCNWQEATSI
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supplemented primate, e.g., human, DTLR4 sequence (SEQ ID NO: 25 and 26); note that nucleotides 81, 3144, 3205, and 3563 designated A, each may be A, C, G, or T; nucleotides 3132, 3532, 3538, and 3553 designated G, each may be G or T; nucleotide 3638 designated A, may be A or T; and nucleotides 3677, 3685, and 3736 designated C, each may be A or C:

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AAAATACTCC CTTGCCTCAA AAACTGCTCG GTCAAACGGT GATAGCAAAC CACGCATTCA       60

CAGGGCCACT GCTGCTCACA AAACCAGTGA GGATGATGCC AGGATG ATG TCT GCC          115
                                                   Met Ser Ala
                                                   -22     -20

TCG CGC CTG GCT GGG ACT CTG ATC CCA GCC ATG GCC TTC CTC TCC TGC         163
Ser Arg Leu Ala Gly Thr Leu Ile Pro Ala Met Ala Phe Leu Ser Cys
               -15                 -10                 -5

GTG AGA CCA GAA AGC TGG GAG CCC TGC GTG GAG GTT CCT AAT ATT ACT         211
Val Arg Pro Glu Ser Trp Glu Pro Cys Val Glu Val Pro Asn Ile Thr
            1               5                  10

TAT CAA TGC ATG GAG CTG AAT TTC TAC AAA ATC CCC GAC AAC CTC CCC         259
Tyr Gln Cys Met Glu Leu Asn Phe Tyr Lys Ile Pro Asp Asn Leu Pro
     15                  20                  25

TTC TCA ACC AAG AAC CTG GAC CTG AGC TTT AAT CCC CTG AGG CAT TTA         307
Phe Ser Thr Lys Asn Leu Asp Leu Ser Phe Asn Pro Leu Arg His Leu
 30                  35                  40                  45

GGC AGC TAT AGC TTC TTC AGT TTC CCA GAA CTG CAG GTG CTC GAT TTA         355
Gly Ser Tyr Ser Phe Phe Ser Phe Pro Glu Leu Gln Val Leu Asp Leu
                50                  55                  60

TCC AGG TGT GAA ATC CAG ACA ATT GAA GAT GGG GCA TAT CAG AGC CTA         403
Ser Arg Cys Glu Ile Gln Thr Ile Glu Asp Gly Ala Tyr Gln Ser Leu
            65                  70                  75

AGC CAC CTC TCT ACC TTA ATA TTG ACA GGA AAC CCC ATC CAG AGT TTA         451
Ser His Leu Ser Thr Leu Ile Leu Thr Gly Asn Pro Ile Gln Ser Leu
         80                  85                  90

GCC CTG GGA GCC TTT TCT GGA CTA TCA AGT TTA CAG AAG CTG GTG GCT         499
Ala Leu Gly Ala Phe Ser Gly Leu Ser Ser Leu Gln Lys Leu Val Ala
     95                  100                 105

GTG GAG ACA AAT CTA GCA TCT CTA GAG AAC TTC CCC ATT GGA CAT CTC         547
Val Glu Thr Asn Leu Ala Ser Leu Glu Asn Phe Pro Ile Gly His Leu
110                 115                 120                 125

AAA ACT TTG AAA GAA CTT AAT GTG GCT CAC AAT CTT ATC CAA TCT TTC         595
Lys Thr Leu Lys Glu Leu Asn Val Ala His Asn Leu Ile Gln Ser Phe
                130                 135                 140

AAA TTA CCT GAG TAT TTT TCT AAT CTG ACC AAT CTA GAG CAC TTG GAC         643
Lys Leu Pro Glu Tyr Phe Ser Asn Leu Thr Asn Leu Glu His Leu Asp
            145                 150                 155
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CTT TCC AGC AAC AAG ATT CAA AGT ATT TAT TGC ACA GAC TTG CGG GTT      691
Leu Ser Ser Asn Lys Ile Gln Ser Ile Tyr Cys Thr Asp Leu Arg Val
        160                 165                 170

CTA CAT CAA ATG CCC CTA CTC AAT CTC TCT TTA GAC CTG TCC CTG AAC      739
Leu His Gln Met Pro Leu Leu Asn Leu Ser Leu Asp Leu Ser Leu Asn
    175                 180                 185

CCT ATG AAC TTT ATC CAA CCA GGT GCA TTT AAA GAA ATT AGG CTT CAT      787
Pro Met Asn Phe Ile Gln Pro Gly Ala Phe Lys Glu Ile Arg Leu His
190                 195                 200                 205

AAG CTG ACT TTA AGA AAT AAT TTT GAT AGT TTA AAT GTA ATG AAA ACT      835
Lys Leu Thr Leu Arg Asn Asn Phe Asp Ser Leu Asn Val Met Lys Thr
                210                 215                 220

TGT ATT CAA GGT CTC CCT GGT TTA GAA GTC CAT CGT TTG GTT CTG GGA      883
Cys Ile Gln Gly Leu Ala Gly Leu Glu Val His Arg Leu Val Leu Gly
            225                 230                 235

GAA TTT AGA AAT GAA GGA AAC TTG GAA AAG TTT GAC AAA TCT GCT CTA      931
Glu Phe Arg Asn Glu Gly Asn Leu Glu Lys Phe Asp Lys Ser Ala Leu
        240                 245                 250

GAG GGC CTG TGC AAT TTG ACC ATT GAA GAA TTC CGA TTA GCA TAC TTA      979
Glu Gly Leu Cys Asn Leu Thr Ile Glu Glu Phe Arg Leu Ala Tyr Leu
    255                 260                 265

GAC TAC TAC CTC GAT GAT ATT ATT GAC TTA TTT AAT TGT TTG ACA AAT     1027
Asp Tyr Tyr Leu Asp Asp Ile Ile Asp Leu Phe Asn Cys Leu Thr Asn
270                 275                 280                 285

GTT TCT TCA TTT TCC CTG GTG AGT GTG ACT ATT GAA AGG GTA AAA GAC     1075
Val Ser Ser Phe Ser Leu Val Ser Val Thr Ile Glu Arg Val Lys Asp
                290                 295                 300

TTT TCT TAT AAT TTC GGA TGG CAA CAT TTA GAA TTA GTT AAC TGT AAA     1123
Phe Ser Tyr Asn Phe Gly Trp Gln His Leu Glu Leu Val Asn Cys Lys
            305                 310                 315

TTT GGA CAG TTT CCC ACA TTG AAA CTC AAA TCT CTC AAA AGG CTT ACT     1171
Phe Gly Gln Phe Pro Thr Leu Lys Leu Lys Ser Leu Lys Arg Leu Thr
        320                 325                 330

TTC ACT TCC AAC AAA GGT GGG AAT GCT TTT TCA GAA GTT GAT CTA CCA     1219
Phe Thr Ser Asn Lys Gly Gly Asn Ala Phe Ser Glu Val Asp Leu Pro
    335                 340                 345

AGC CTT GAG TTT CTA GAT CTC AGT AGA AAT GGC TTG AGT TTC AAA GGT     1267
Ser Leu Glu Phe Leu Asp Leu Ser Arg Asn Gly Leu Ser Phe Lys Gly
350                 355                 360                 365
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TGC TGT TCT CAA AGT GAT TTT GGG ACA ACC AGC CTA AAG TAT TTA GAT      1315
Cys Cys Ser Gln Ser Asp Phe Gly Thr Thr Ser Leu Lys Tyr Leu Asp
                370                 375                 380

CTG AGC TTC AAT GGT GTT ATT ACC ATG AGT TCA AAC TTC TTG GGC TTA      1363
Leu Ser Phe Asn Gly Val Ile Thr Met Ser Ser Asn Phe Leu Gly Leu
            385                 390                 395

GAA CAA CTA GAA CAT CTG GAT TTC CAG CAT TCC AAT TTG AAA CAA ATG      1411
Glu Gln Leu Glu His Leu Asp Phe Gln His Ser Asn Leu Lys Gln Met
        400                 405                 410

AGT GAG TTT TCA GTA TTC CTA TCA CTC AGA AAC CTC ATT TAC CTT GAC      1459
Ser Glu Phe Ser Val Phe Leu Ser Leu Arg Asn Leu Ile Tyr Leu Asp
    415                 420                 425

ATT TCT CAT ACT CAC ACC AGA GTT GCT TTC AAT GGC ATC TTC AAT GGC      1507
Ile Ser His Thr His Thr Arg Val Ala Phe Asn Gly Ile Phe Asn Gly
430                 435                 440                 445

TTG TCC AGT CTC GAA GTC TTG AAA ATG GCT GGC AAT TCT TTC CAG GAA      1555
Leu Ser Ser Leu Glu Val Leu Lys Met Ala Gly Asn Ser Phe Gln Glu
                450                 455                 460

AAC TTC CTT CCA GAT ATC TTC ACA GAG CTG AGA AAC TTG ACC TTC CTG      1603
Asn Phe Leu Pro Asp Ile Phe Thr Glu Leu Arg Asn Leu Thr Phe Leu
            465                 470                 475

GAC CTC TCT CAG TGT CAA CTG GAG CAG TTG TCT CCA ACA GCA TTT AAC      1651
Asp Leu Ser Gln Cys Gln Leu Glu Gln Leu Ser Pro Thr Ala Phe Asn
        480                 485                 490

TCA CTC TCC AGT CTT CAG GTA CTA AAT ATG AGC CAC AAC AAC TTC TTT      1699
Ser Leu Ser Ser Leu Gln Val Leu Asn Met Ser His Asn Asn Phe Phe
    495                 500                 505

TCA TTG GAT ACG TTT CCT TAT AAG TGT CTG AAC TCC CTC CAG GTT CTT      1747
Ser Leu Asp Thr Phe Pro Tyr Lys Cys Leu Asn Ser Leu Gln Val Leu
510                 515                 520                 525

GAT TAC AGT CTC AAT CAC ATA ATG ACT TCC AAA AAA CAG GAA CTA CAG      1795
Asp Tyr Ser Leu Asn His Ile Met Thr Ser Lys Lys Gln Glu Leu Gln
                530                 535                 540

CAT TTT CCA AGT AGT CTA GCT TTC TTA AAT CTT ACT CAG AAT GAC TTT      1843
His Phe Pro Ser Ser Leu Ala Phe Leu Asn Leu Thr Gln Asn Asp Phe
            545                 550                 555

GCT TGT ACT TGT GAA CAC CAG AGT TTC CTG CAA TGG ATC AAG GAC CAG      1891
Ala Cys Thr Cys Glu His Gln Ser Phe Leu Gln Trp Ile Lys Asp Gln
        560                 565                 570

AGG CAG CTC TTG GTG GAA GTT GAA CGA ATG GAA TGT GCA ACA CCT TCA      1939
Arg Gln Leu Leu Val Glu Val Glu Arg Met Glu Cys Ala Thr Pro Ser
    575                 580                 585
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GAT AAG CAG GGC ATG CCT GTG CTG AGT TTG AAT ATC ACC TGT CAG ATG      1987
Asp Lys Gln Gly Met Pro Val Leu Ser Leu Asn Ile Thr Cys Gln Met
590                 595                 600                 605

AAT AAG ACC ATC ATT GGT GTG TCG GTC CTC AGT GTG CTT GTA GTA TCT      2035
Asn Lys Thr Ile Ile Gly Val Ser Val Leu Ser Val Leu Val Val Ser
                610                 615                 620

GTT GTA GCA GTT CTG GTC TAT AAG TTC TAT TTT CAC CTG ATG CTT CTT      2083
Val Val Ala Val Leu Val Tyr Lys Phe Tyr Phe His Leu Met Leu Leu
            625                 630                 635

GCT GGC TGC ATA AAG TAT GGT AGA GGT GAA AAC ATC TAT GAT GCC TTT      2131
Ala Gly Cys Ile Lys Tyr Gly Arg Gly Glu Asn Ile Tyr Asp Ala Phe
        640                 645                 650

GTT ATC TAC TCA AGC CAG GAT GAG GAC TGG GTA AGG AAT GAG CTA GTA      2179
Val Ile Tyr Ser Ser Gln Asp Glu Asp Trp Val Arg Asn Glu Leu Val
    655                 660                 665

AAG AAT TTA GAA GAA GGG GTG CCT CCA TTT CAG CTC TGC CTT CAC TAC      2227
Lys Asn Leu Glu Glu Gly Val Pro Pro Phe Gln Leu Cys Leu His Tyr
670                 675                 680                 685

AGA GAC TTT ATT CCC GGT GTG GCC ATT GCT GCC AAC ATC ATC CAT GAA      2275
Arg Asp Phe Ile Pro Gly Val Ala Ile Ala Ala Asn Ile Ile His Glu
                690                 695                 700

GGT TTC CAT AAA AGC CGA AAG GTG ATT GTT GTG GTG TCC CAG CAC TTC      2323
Gly Phe His Lys Ser Arg Lys Val Ile Val Val Val Ser Gln His Phe
            705                 710                 715

ATC CAG AGC CGC TGG TGT ATC TTT GAA TAT GAG ATT GCT CAG ACC TGG      2371
Ile Gln Ser Arg Trp Cys Ile Phe Glu Tyr Glu Ile Ala Gln Thr Trp
        720                 725                 730

CAG TTT CTG AGC AGT CGT GCT GGT ATC ATC TTC ATT GTC CTG CAG AAG      2419
Gln Phe Leu Ser Ser Arg Ala Gly Ile Ile Phe Ile Val Leu Gln Lys
    735                 740                 745

GTG GAG AAG ACC CTG CTC AGG CAG CAG GTG GAG CTG TAC CGC CTT CTC      2467
Val Glu Lys Thr Leu Leu Arg Gln Gln Val Glu Leu Tyr Arg Leu Leu
750                 755                 760                 765

AGC AGG AAC ACT TAC CTG GAG TGG GAG GAC AGT GTC CTG GGG CGG CAC      2515
Ser Arg Asn Thr Tyr Leu Glu Trp Glu Asp Ser Val Leu Gly Arg His
                770                 775                 780

ATC TTC TGG AGA CGA CTC AGA AAA GCC CTG CTG GAT GGT AAA TCA TGG      2563
Ile Phe Trp Arg Arg Leu Arg Lys Ala Leu Leu Asp Gly Lys Ser Trp
            785                 790                 795
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AAT CCA GAA GGA ACA GTG GGT ACA GGA TGC AAT TGG CAG GAA GCA ACA      2611
Asn Pro Glu Gly Thr Val Gly Thr Gly Cys Asn Trp Gln Glu Ala Thr
        800                 805                 810

TCT ATC TGAAGAGGAA AAATAAAAAC CTCCTGAGGC ATTTCTTGCC CAGCTGGGTC      2667
Ser Ile
    815

CAACACTTGT TCAGTTAATA AGTATTAAAT GCTGCCACAT GTCAGGCCTT ATGCTAAGGG   2727

TGAGTAATTC CATGGTGCAC TAGATATGCA GGGCTGCTAA TCTCAAGGAG CTTCCAGTGC   2787

AGAGGGAATA AATGCTAGAC TAAAATACAG AGTCTTCCAG GTGGGCATTT CAACCAACTC   2847

AGTCAAGGAA CCCATGACAA AGAAAGTCAT TTCAACTCTT ACCTCATCAA GTTGAATAAA   2907

GACAGAGAAA ACAGAAAGAG ACATTGTTCT TTTCCTGAGT CTTTTGAATG GAAATTGTAT   2967

TATGTTATAG CCATCATAAA ACCATTTTGG TAGTTTTGAC TGAACTGGGT GTTCACTTTT   3027

TCCTTTTTGA TTGAATACAA TTTAAATTCT ACTTGATGAC TGCAGTCGTC AAGGGGCTCC   3087

TGATGCAAGA TGCCCCTTCC ATTTTAAGTC TGTCTCCTTA CAGAGGTTAA AGTCTAATGG   3147

CTAATTCCTA AGGAAACCTG ATTAACACAT GCTCACAACC ATCCTGGTCA TTCTCGAACA   3207

TGTTCTATTT TTTAACTAAT CACCCCTGAT ATATTTTTAT TTTTATATAT CCAGTTTTCA   3267

TTTTTTTACG TCTTGCCTAT AAGCTAATAT CATAAATAAG GTTGTTTAAG ACGTGCTTCA   3327

AATATCCATA TTAACCACTA TTTTTCAAGG AAGTATGGAA AAGTACACTC TGTCACTTTG   3387

TCACTCGATG TCATTCCAAA GTTATTGCCT ACTAAGTAAT GACTGTCATG AAAGCAGCAT   3447

TGAAATAATT TGTTTAAAGG GGGCACTCTT TTAAACGGGA AGAAAATTTC CGCTTCCTGG   3507

TCTTATCATG GACAATTTGG GCTAGAGGCA GGAAGGAAGT GGGATGACCT CAGGAAGTCA   3567

CCTTTTCTTG ATTCCAGAAA CATATGGGCT GATAAACCCG GGGTGACCTC ATGAAATGAG   3627

TTGCAGCAGA AGTTTATTTT TTTCAGAACA AGTGATGTTT GATGGACCTC TGAATCTCTT   3687

TAGGGAGACA CAGATGGCTG GGATCCCTCC CCTGTACCCT TCTCACTGCC AGGAGAACTA   3747

CGTGTGAAGG TATTCAACGC AGGGAGTATA CATTGCTGTT TCCTGTTGGG CAATGCTCCT   3807

TGACCACATT TTGGGAAGAG TGGATGTTAT CATTGAGAAA ACAATGTGTC TGGAATTAAT   3867

GGGGTTCTTA TAAAGAAGGT TCCCAGAAAA GAATGTTCAT TCCAGCTTCT TCAGGAAACA   3927

GGAACATTCA AGGAAAAGGA CAATCAGGAT GTCATCAGGG AAATGAAAAT AAAAACCACA   3987

ATGAGATATC ACCTTATACC AGGTAGATGG CTACTATAAA AAAATGAAGT GTCATCAAGG   4047

ATATAGAGAA ATTGGAACCC TTCTTCACTG CTGGAGGGAA TGGAAAATGG TGTAGCCGTT   4107
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ATCAAAAACA GTACGGAGGT TTCTCAAAAA TTAAAAATAG AACTGCTATA TGATCCAGCA 4167

ATCTCACTTC TGTATATATA CCCAAAATAA TTGAAATCAG AATTTCAAGA AAATATTTAC 4227

ACTCCCATGT TCATTGTGGC ACTCTTCACA ATCACTGTTT CCAAAGTTAT GGAAACAACC 4287

CAAATTTCCA TTGGAAAATA AATGGACAAA GGAAATGTGC ATATAACGTA CAATGGGGAT 4347

ATTATTCAGC CTAAAAAAAG GGGGGATCCT GTTATTTATG ACAACATGAA TAAACCCGGA 4407

GGCCATTATG CTATGTAAAA TGAGCAAGTA ACAGAAAGAC AAATACTGCC TGATTTCATT 4467

TATATGAGGT TCTAAAATAG TCAAACTCAT AGAAGCAGAG AATAGAACAG TGGTTCCTAG 4527

GGAAAAGGAG GAAGGGAGAA ATGAGGAAAT AGGGAGTTGT CTAATTGGTA TAAAATTATA 4587

GTATGCAAGA TGAATTAGCT CTAAAGATCA GCTGTATAGC AGAGTTCGTA TAATGAACAA 4647

TACTGTATTA TGCACTTAAC ATTTTGTTAA GAGGGTACCT CTCATGTTAA GTGTTCTTAC 4707

CATATACATA TACACAAGGA AGCTTTTGGA GGTGATGGAT ATATTTATTA CCTTGATTGT 4767

GGTGATGGTT TGACAGGTAT GTGACTATGT CTAAACTCAT CAAATTGTAT ACATTAAATA 4827

TATGCAGTTT TATAATATCA AAAAAAAAAA AAAAAAAA 4865

MSASRLAGTLIPAMAFLSCVRPESWEPCVEVPNITYQCMELNFYKIPDNLPFSTKNLDLSFNPLRHLGSYSFFSF
PELQVLDLSRCEIQTIEDGAYQSLSHLSTLILTGNPIQSLALGAFSGLSSLQKLVAVETNLASLENFPIGHLKTL
KELNVAHNLIQSFKLPEYFSNLTNLEHLDLSSNKIQSIYCTDLRVLHQMPLLNLSLDLSLNPMNFIQPGAFKEIR
LHKLTLRNNFDSLNVMKTCIQGLAGLEVHRLVLGEFRNEGNLEKFDKSALEGLCNLTIEEFRLAYLDYYLDDIID
LFNCLTNVSSFSLVSVTIERVKDFSYNFGWQHLELVNCKFGQFPTLKLKSLKRLTFTSNKGGNAFSEVDLPSLEF
LDLSRNGLSFKGCCSQSDFGTTSLKYLDLSFNGVITMSSNFLGLEQLEHLDFQHSNLKQMSEFSVFLSLRNLIYL
DISHTHTRVAFNGIFNGLSSLEVLKMAGNSFQENFLPDIFTELRNLTFLDLSQCQLEQLSPTAFNSLSSLQVLNM
SHNNFFSLDTFPYKCLNSLQVLDYSLNHIMTSKKQELQHFPSSLAFLNLTQNDFACTCEHQSFLQWIKDQRQLLV
EVERMECATPSDKQGMPVLSLNITCQMNKTIIGVSVLSVLVVSVVAVLVYKFYFHLMLLAGCIKYGRGENIYDAF
VIYSSQDEDWVRNELVKNLEEGVPPFQLCLHYRDFIPGVAIAANIIHEGFHKSRKVIVVVSQHFIQSRWCIFEYE
IAQTWQFLSSRAGIIFIVLQKVEKTLLRQQVELYRLLSRNTYLEWEDSVLGRHIFWRRLRKALLDGKSWNPEGTV
GTGCNWQEATSI

Table 5: Partial nucleotide and amino acid sequences (see SEQ ID NO: 9 and 10) of a mammalian, e.g., primate, human, DNAX Toll like Receptor 5 (DTLR5).

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TGT TGG GAT GTC TTT GAG GGA CTT TCT CAT CTT CAA GTT CTG TAT TTG          48
Cys Trp Asp Val Phe Glu Gly Leu Ser His Leu Gln Val Leu Tyr Leu
  1               5                  10                  15

AAT CAT AAC TAT CTT AAT TCC CTT CCA CCA GGA GTA TTT AGC CAT CTG          96
Asn His Asn Tyr Leu Asn Ser Leu Pro Pro Gly Val Phe Ser His Leu
             20                  25                  30

ACT GCA TTA AGG GGA CTA AGC CTC AAC TCC AAC AGG CTG ACA GTT CTT         144
Thr Ala Leu Arg Gly Leu Ser Leu Asn Ser Asn Arg Leu Thr Val Leu
         35                  40                  45

TCT CAC AAT GAT TTA CCT GCT AAT TTA GAG ATC CTG GAC ATA TCC AGG         192
Ser His Asn Asp Leu Pro Ala Asn Leu Glu Ile Leu Asp Ile Ser Arg
     50                  55                  60

AAC CAG CTC CTA GCT CCT AAT CCT GAT GTA TTT GTA TCA CTT AGT GTC         240
Asn Gln Leu Leu Ala Pro Asn Pro Asp Val Phe Val Ser Leu Ser Val
 65                  70                  75                  80

TTG GAT ATA ACT CAT AAC AAG TTC ATT TGT GAA TGT GAA CTT AGC ACT         288
Leu Asp Ile Thr His Asn Lys Phe Ile Cys Glu Cys Glu Leu Ser Thr
                 85                  90                  95

TTT ATC AAT TGG CTC AAT CAC ACC AAT GTC ACT ATA GCT GGG CCT CCT         336
Phe Ile Asn Trp Leu Asn His Thr Asn Val Thr Ile Ala Gly Pro Pro
            100                 105                 110

GCA GAC ATA TAT TGT GTG TAC CCT GAC TCG TTC TCT GGG GTT TCC CTC         384
Ala Asp Ile Tyr Cys Val Tyr Pro Asp Ser Phe Ser Gly Val Ser Leu
        115                 120                 125

TTC TCT CTT TCC ACG GAA GGT TGT GAT GAA GAG GAA GTC TTA AAG TCC         432
Phe Ser Leu Ser Thr Glu Gly Cys Asp Glu Glu Glu Val Leu Lys Ser
    130                 135                 140

CTA AAG TTC TCC CTT TTC ATT GTA TGC ACT GTC ACT CTG ACT CTG TTC         480
Leu Lys Phe Ser Leu Phe Ile Val Cys Thr Val Thr Leu Thr Leu Phe
145                 150                 155                 160

CTC ATG ACC ATC CTC ACA GTC ACA AAG TTC CGG GGC TTC TGT TTT ATC         528
Leu Met Thr Ile Leu Thr Val Thr Lys Phe Arg Gly Phe Cys Phe Ile
                165                 170                 175

TGT TAT AAG ACA GCC CAG AGA CTG GTG TTC AAG GAC CAT CCC CAG GGC         576
Cys Tyr Lys Thr Ala Gln Arg Leu Val Phe Lys Asp His Pro Gln Gly
            180                 185                 190

ACA GAA CCT GAT ATG TAC AAA TAT GAT GCC TAT TTG TGC TTC AGC AGC         624
Thr Glu Pro Asp Met Tyr Lys Tyr Asp Ala Tyr Leu Cys Phe Ser Ser
        195                 200                 205
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AAA GAC TTC ACA TGG GTG CAG AAT GCT TTG CTC AAA CAC CTG GAC ACT      672
Lys Asp Phe Thr Trp Val Gln Asn Ala Leu Leu Lys His Leu Asp Thr
    210                 215                 220

CAA TAC AGT GAC CAA AAC AGA TTC AAC CTG TGC TTT GAA GAA AGA GAC      720
Gln Tyr Ser Asp Gln Asn Arg Phe Asn Leu Cys Phe Glu Glu Arg Asp
225                 230                 235                 240

TTT GTC CCA GGA GAA AAC CGC ATT GCC AAT ATC CAG GAT GCC ATC TGG      768
Phe Val Pro Gly Glu Asn Arg Ile Ala Asn Ile Gln Asp Ala Ile Trp
                245                 250                 255

AAC AGT AGA AAG ATC GTT TGT CTT GTG AGC AGA CAC TTC CTT AGA GAT      816
Asn Ser Arg Lys Ile Val Cys Leu Val Ser Arg His Phe Leu Arg Asp
            260                 265                 270

GGC TGG TGC CTT GAA GCC TTC AGT TAT GCC CAG GGC AGG TGC CTA TCT      864
Gly Trp Cys Leu Glu Ala Phe Ser Tyr Ala Gln Gly Arg Cys Leu Ser
        275                 280                 285

GAC CTT AAC AGT GCT CTC ATC ATG GTG GTG GTT GGG TCC TTG TCC CAG      912
Asp Leu Asn Ser Ala Leu Ile Met Val Val Val Gly Ser Leu Ser Gln
    290                 295                 300

TAC CAG TTG ATG AAA CAT CAA TCC ATC AGA GGC TTT GTA CAG AAA CAG      960
Tyr Gln Leu Met Lys His Gln Ser Ile Arg Gly Phe Val Gln Lys Gln
305                 310                 315                 320

CAG TAT TTG AGG TGG CCT GAG GAT CTC CAG GAT GTT GGC TGG TTT CTT     1008
Gln Tyr Leu Arg Trp Pro Glu Asp Leu Gln Asp Val Gly Trp Phe Leu
                325                 330                 335

CAT AAA CTC TCT CAA CAG ATA CTA AAG AAA GAA AAG GAA AAG AAG AAA     1056
His Lys Leu Ser Gln Gln Ile Leu Lys Lys Glu Lys Glu Lys Lys Lys
            340                 345                 350

GAC AAT AAC ATT CCG TTG CAA ACT GTA GCA ACC ATC TCC TAATCAAAGG      1105
Asp Asn Asn Ile Pro Leu Gln Thr Val Ala Thr Ile Ser
        355                 360                 365

AGCAATTTCC AACTTATCTC AAGCCACAAA TAACTCTTCA CTTTGTATTT GCACCAAGTT   1165

ATCATTTTGG GGTCCTCTCT GGAGGCTTCT TTTTTCTTTT TGCTACTATG AAAACAACAT   1225

AAATCTCTCA ATTTTCGTAT CAAAAAAAAA AAAAAAAAAA TGGCGGCCGC              1275

CWDVFEGLSHLQVLYLNHNYLNSLPPGVFSHLTALRGLSLNSNRLTVLSHNDLPANLEILDISRNQLLAPNPDVF
VSLSVLDITHNKFICECELSTFINWLNHTNVTIAGPPADIYCVYPDSFSGVSLFSLSTEGCDEEEVLKSLKFSLF
IVCTVTLTLFLMTILTVTKFRGFCFICYKTAQRLVFKDHPQGTEPDMYKYDAYLCFSSKDFTWVQNALLKHLDTQ
YSDQNRFNLCFEERDFVPGENRIANIQDAIWNSRKIVCLVSRHFLRDGWCLEAFSYAQGRCLSDLNSALIMVVVG
SLSQYQLMKHQSIRGFVQKQQYLRWPEDLQDVGWFLHKLSQQILKKEKEKKKDNNIPLQTVATIS
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Table 6: Nucleotide and amino acid sequences of mammalian, e.g., primate or rodent DNAX Toll like Receptor 6 (DTLR6). SEQ ID NO: 11 and 12 are from primate, e.g., human; SEQ ID NO: 13 and 14 are from rodent, e.g., mouse.

primate:

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ATG TGG ACA CTG AAG AGA CTA ATT CTT ATC CCT TTT AAC ATA ATC CTA          48
Met Trp Thr Leu Lys Arg Leu Ile Leu Ile Leu Phe Asn Ile Ile Leu
-22  -20                 -15                 -10

ATT TCC AAA CTC CTT GGG GCT AGA TGG TTT CCT AAA ACT CTG CCC TGT          96
Ile Ser Lys Leu Leu Gly Ala Arg Trp Phe Pro Lys Thr Leu Pro Cys
 -5                   1                   5                  10

GAT GTC ACT CTG GAT GTT CCA AAG AAC CAT GTG ATC GTG GAC TGC ACA         144
Asp Val Thr Leu Asp Val Pro Lys Asn His Val Ile Val Asp Cys Thr
                15                  20                  25

GAC AAG CAT TTG ACA GAA ATT CCT GGA GGT ATT CCC ACG AAC ACC ACG         192
Asp Lys His Leu Thr Glu Ile Pro Gly Gly Ile Pro Thr Asn Thr Thr
            30                  35                  40

AAC CTC ACC CTC ACC ATT AAC CAC ATA CCA GAC ATC TCC CCA GCG TCC         240
Asn Leu Thr Leu Thr Ile Asn His Ile Pro Asp Ile Ser Pro Ala Ser
        45                  50                  55

TTT CAC AGA CTC GAC CAT CTG GTA GAG ATC GAT TTC AGA TGC AAC TGT         288
Phe His Arg Leu Asp His Leu Val Glu Ile Asp Phe Arg Cys Asn Cys
    60                  65                  70

GTA CCT ATT CCA CTG GGG TCA AAA AAC AAC ATG TGC ATC AAG AGG CTG         336
Val Pro Ile Pro Leu Gly Ser Lys Asn Asn Met Cys Ile Lys Arg Leu
75                  80                  85                  90

CAG ATT AAA CCC AGA AGC TTT AGT GGA CTC ACT TAC TTA AAA TCC CTT         384
Gln Ile Lys Pro Arg Ser Phe Ser Gly Leu Thr Tyr Leu Lys Ser Leu
                95                 100                 105

TAC CTG GAT GGA AAC CAG CTA CTA GAG ATA CCG CAG GGC CTC CCG CCT         432
Tyr Leu Asp Gly Asn Gln Leu Leu Glu Ile Pro Gln Gly Leu Pro Pro
            110                 115                 120

AGC TTA CAG CTT CTC AGC CTT GAG GCC AAC AAC ATC TTT TCC ATC AGA         480
Ser Leu Gln Leu Leu Ser Leu Glu Ala Asn Asn Ile Phe Ser Ile Arg
        125                 130                 135

AAA GAG AAT CTA ACA GAA CTG GCC AAC ATA GAA ATA CTC TAC CTG GGC         528
Lys Glu Asn Leu Thr Glu Leu Ala Asn Ile Glu Ile Leu Tyr Leu Gly
    140                 145                 150

CAA AAC TGT TAT TAT CGA AAT CCT TGT TAT GTT TCA TAT TCA ATA GAG         576
Gln Asn Cys Tyr Tyr Arg Asn Pro Cys Tyr Val Ser Tyr Ser Ile Glu
155                 160                 165                 170
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AAA GAT GCC TTC CTA AAC TTG ACA AAG TTA AAA GTG CTC TCC CTG AAA      624
Lys Asp Ala Phe Leu Asn Leu Thr Lys Leu Lys Val Leu Ser Leu Lys
                175                 180                 185

GAT AAC AAT GTC ACA GCC GTC CCT ACT GTT TTG CCA TCT ACT TTA ACA      672
Asp Asn Asn Val Thr Ala Val Pro Thr Val Leu Pro Ser Thr Leu Thr
            190                 195                 200

GAA CTA TAT CTC TAC AAC AAC ATG ATT GCA AAA ATC CAA GAA GAT GAT      720
Glu Leu Tyr Leu Tyr Asn Asn Met Ile Ala Lys Ile Gln Glu Asp Asp
        205                 210                 215

TTT AAT AAC CTC AAC CAA TTA CAA ATT CTT GAC CTA AGT GGA AAT TGC      768
Phe Asn Asn Leu Asn Gln Leu Gln Ile Leu Asp Leu Ser Gly Asn Cys
    220                 225                 230

CCT CGT TGT TAT AAT GCC CCA TTT CCT TGT GCG CCG TGT AAA AAT AAT      816
Pro Arg Cys Tyr Asn Ala Pro Phe Pro Cys Ala Pro Cys Lys Asn Asn
235                 240                 245                 250

TCT CCC CTA CAG ATC CCT GTA AAT GCT TTT GAT GCG CTG ACA GAA TTA      864
Ser Pro Leu Gln Ile Pro Val Asn Ala Phe Asp Ala Leu Thr Glu Leu
                255                 260                 265

AAA GTT TTA CGT CTA CAC AGT AAC TCT CTT CAG CAT GTG CCC CCA AGA      912
Lys Val Leu Arg Leu His Ser Asn Ser Leu Gln His Val Pro Pro Arg
            270                 275                 280

TGG TTT AAG AAC ATC AAC AAA CTC CAG GAA CTG GAT CTG TCC CAA AAC      960
Trp Phe Lys Asn Ile Asn Lys Leu Gln Glu Leu Asp Leu Ser Gln Asn
        285                 290                 295

TTC TTG GCC AAA GAA ATT GGG GAT GCT AAA TTT CTG CAT TTT CTC CCC     1008
Phe Leu Ala Lys Glu Ile Gly Asp Ala Lys Phe Leu His Phe Leu Pro
    300                 305                 310

AGC CTC ATC CAA TTG GAT CTG TCT TTC AAT TTT GAA CTT CAG GTC TAT     1056
Ser Leu Ile Gln Leu Asp Leu Ser Phe Asn Phe Glu Leu Gln Val Tyr
315                 320                 325                 330

CGT GCA TCT ATG AAT CTA TCA CAA GCA TTT TCT TCA CTG AAA AGC CTG     1104
Arg Ala Ser Met Asn Leu Ser Gln Ala Phe Ser Ser Leu Lys Ser Leu
                335                 340                 345

AAA ATT CTG CGG ATC AGA GGA TAT GTC TTT AAA GAG TTG AAA AGC TTT     1152
Lys Ile Leu Arg Ile Arg Gly Tyr Val Phe Lys Glu Leu Lys Ser Phe
            350                 355                 360

AAC CTC TCG CCA TTA CAT AAT CTT CAA AAT CTT GAA GTT CTT GAT CTT     1200
Asn Leu Ser Pro Leu His Asn Leu Gln Asn Leu Glu Val Leu Asp Leu
        365                 370                 375

GGC ACT AAC TTT ATA AAA ATT GCT AAC CTC AGC ATG TTT AAA CAA TTT     1248
Gly Thr Asn Phe Ile Lys Ile Ala Asn Leu Ser Met Phe Lys Gln Phe
    380                 385                 390
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AAA AGA CTG AAA GTC ATA GAT CTT TCA GTG AAT AAA ATA TCA CCT TCA      1296
Lys Arg Leu Lys Val Ile Asp Leu Ser Val Asn Lys Ile Ser Pro Ser
395                 400                 405                 410

GGA GAT TCA AGT GAA GTT GGC TTC TGC TCA AAT GCC AGA ACT TCT GTA      1344
Gly Asp Ser Ser Glu Val Gly Phe Cys Ser Asn Ala Arg Thr Ser Val
                415                 420                 425

GAA AGT TAT GAA CCC CAG GTC CTG GAA CAA TTA CAT TAT TTC AGA TAT      1392
Glu Ser Tyr Glu Pro Gln Val Leu Glu Gln Leu His Tyr Phe Arg Tyr
            430                 435                 440

GAT AAG TAT GCA AGG AGT TGC AGA TTC AAA AAC AAA GAG GCT TCT TTC      1440
Asp Lys Tyr Ala Arg Ser Cys Arg Phe Lys Asn Lys Glu Ala Ser Phe
        445                 450                 455

ATG TCT GTT AAT GAA AGC TGC TAC AAG TAT GGG CAG ACC TTG GAT CTA      1488
Met Ser Val Asn Glu Ser Cys Tyr Lys Tyr Gly Gln Thr Leu Asp Leu
    460                 465                 470

AGT AAA AAT AGT ATA TTT TTT GTC AAG TCC TCT GAT TTT CAG CAT CTT      1536
Ser Lys Asn Ser Ile Phe Phe Val Lys Ser Ser Asp Phe Gln His Leu
475                 480                 485                 490

TCT TTC CTC AAA TGC CTG AAT CTG TCA GGA AAT CTC ATT AGC CAA ACT      1584
Ser Phe Leu Lys Cys Leu Asn Leu Ser Gly Asn Leu Ile Ser Gln Thr
                495                 500                 505

CTT AAT GGC AGT GAA TTC CAA CCT TTA GCA GAG CTG AGA TAT TTG GAC      1632
Leu Asn Gly Ser Glu Phe Gln Pro Leu Ala Glu Leu Arg Tyr Leu Asp
            510                 515                 520

TTC TCC AAC AAC CGG CTT GAT TTA CTC CAT TCA ACA GCA TTT GAA GAG      1680
Phe Ser Asn Asn Arg Leu Asp Leu Leu His Ser Thr Ala Phe Glu Glu
        525                 530                 535

CTT CAC AAA CTG GAA GTT CTG GAT ATA AGC AGT AAT AGC CAT TAT TTT      1728
Leu His Lys Leu Glu Val Leu Asp Ile Ser Ser Asn Ser His Tyr Phe
    540                 545                 550

CAA TCA GAA GGA ATT ACT CAT ATG CTA AAC TTT ACC AAG AAC CTA AAG      1776
Gln Ser Glu Gly Ile Thr His Met Leu Asn Phe Thr Lys Asn Leu Lys
555                 560                 565                 570

GTT CTG CAG AAA CTG ATG ATG AAC GAC AAT GAC ATC TCT TCC TCC ACC      1824
Val Leu Gln Lys Leu Met Met Asn Asp Asn Asp Ile Ser Ser Ser Thr
                575                 580                 585

AGC AGG ACC ATG GAG AGT GAG TCT CTT AGA ACT CTG GAA TTC AGA GGA      1872
Ser Arg Thr Met Glu Ser Glu Ser Leu Arg Thr Leu Glu Phe Arg Gly
            590                 595                 600
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AAT CAC TTA GAT GTT TTA TGG AGA GAA GGT GAT AAC AGA TAC TTA CAA      1920
Asn His Leu Asp Val Leu Trp Arg Glu Gly Asp Asn Arg Tyr Leu Gln
        605                 610                 615

TTA TTC AAG AAT CTG CTA AAA TTA GAG GAA TTA GAC ATC TCT AAA AAT      1968
Leu Phe Lys Asn Leu Leu Lys Leu Glu Glu Leu Asp Ile Ser Lys Asn
    620                 625                 630

TCC CTA AGT TTC TTG CCT TCT GGA GTT TTT GAT GGT ATG CCT CCA AAT      2016
Ser Leu Ser Phe Leu Pro Ser Gly Val Phe Asp Gly Met Pro Pro Asn
635                 640                 645                 650

CTA AAG AAT CTC TCT TTG GCC AAA AAT GGG CTC AAA TCT TTC AGT TGG      2064
Leu Lys Asn Leu Ser Leu Ala Lys Asn Gly Leu Lys Ser Phe Ser Trp
                655                 660                 665

AAG AAA CTC CAG TGT CTA AAG AAC CTG GAA ACT TTG GAC CTC AGC CAC      2112
Lys Lys Leu Gln Cys Leu Lys Asn Leu Glu Thr Leu Asp Leu Ser His
            670                 675                 680

AAC CAA CTG ACC ACT GTC CCT GAG AGA TTA TCC AAC TGT TCC AGA AGC      2160
Asn Gln Leu Thr Thr Val Pro Glu Arg Leu Ser Asn Cys Ser Arg Ser
        685                 690                 695

CTC AAG AAT CTG ATT CTT AAG AAT AAT CAA ATC AGG AGT CTG ACG AAG      2208
Leu Lys Asn Leu Ile Leu Lys Asn Asn Gln Ile Arg Ser Leu Thr Lys
    700                 705                 710

TAT TTT CTA CAA GAT GCC TTC CAG TTG CGA TAT CTG GAT CTC AGC TCA      2256
Tyr Phe Leu Gln Asp Ala Phe Gln Leu Arg Tyr Leu Asp Leu Ser Ser
715                 720                 725                 730

AAT AAA ATC CAG ATG ATC CAA AAG ACC AGC TTC CCA GAA AAT GTC CTC      2304
Asn Lys Ile Gln Met Ile Gln Lys Thr Ser Phe Pro Glu Asn Val Leu
                735                 740                 745

AAC AAT CTG AAG ATG TTG CTT TTG CAT CAT AAT CGG TTT CTG TGC ACC      2352
Asn Asn Leu Lys Met Leu Leu Leu His His Asn Arg Phe Leu Cys Thr
            750                 755                 760

TGT GAT GCT GTG TGG TTT GTC TGG TGG GTT AAC CAT ACG GAG GTG ACT      2400
Cys Asp Ala Val Trp Phe Val Trp Trp Val Asn His Thr Glu Val Thr
        765                 770                 775

ATT CCT TAC CTG GCC ACA GAT GTG ACT TGT GTG GGG CCA GGA GCA CAC      2448
Ile Pro Tyr Leu Ala Thr Asp Val Thr Cys Val Gly Pro Gly Ala His
    780                 785                 790

AAG GGC CAA AGT GTG ATC TCC CTG GAT CTG TAC ACC TGT GAG TTA GAT      2496
Lys Gly Gln Ser Val Ile Ser Leu Asp Leu Tyr Thr Cys Glu Leu Asp
795                 800                 805                 810

CTG ACT AAC CTG ATT CTG TTC TCA CTT TCC ATA TCT GTA TCT CTC TTT      2544
Leu Thr Asn Leu Ile Leu Phe Ser Leu Ser Ile Ser Val Ser Leu Phe
                815                 820                 825
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CTC ATG GTG ATG ATG ACA GCA AGT CAC CTC TAC TTC TGG GAT GTG TGG      2592
Leu Met Val Met Met Thr Ala Ser His Leu Tyr Phe Trp Asp Val Trp
            830                 835                 840

TAT ATT TAC CAT TTC TGT AAG GCC AAG ATA AAG GGG TAT CAG CGT CTA      2640
Tyr Ile Tyr His Phe Cys Lys Ala Lys Ile Lys Gly Tyr Gln Arg Leu
        845                 850                 855

ATA TCA CCA GAC TGT TGC TAT GAT GCT TTT ATT GTG TAT GAC ACT AAA      2688
Ile Ser Pro Asp Cys Cys Tyr Asp Ala Phe Ile Val Tyr Asp Thr Lys
    860                 865                 870

GAC CCA GCT GTG ACC GAG TGG GTT TTG GCT GAG CTG GTG GCC AAA CTG      2736
Asp Pro Ala Val Thr Glu Trp Val Leu Ala Glu Leu Val Ala Lys Leu
875                 880                 885                 890

GAA GAC CCA AGA GAG AAA CAT TTT AAT TTA TGT CTC GAG GAA AGG GAC      2784
Glu Asp Pro Arg Glu Lys His Phe Asn Leu Cys Leu Glu Glu Arg Asp
                895                 900                 905

TGG TTA CCA GGG CAG CCA GTT CTG GAA AAC CTT TCC CAG AGC ATA CAG      2832
Trp Leu Pro Gly Gln Pro Val Leu Glu Asn Leu Ser Gln Ser Ile Gln
            910                 915                 920

CTT AGC AAA AAG ACA GTG TTT GTG ATG ACA GAC AAG TAT GCA AAG ACT      2880
Leu Ser Lys Lys Thr Val Phe Val Met Thr Asp Lys Tyr Ala Lys Thr
        925                 930                 935

GAA AAT TTT AAG ATA GCA TTT TAC TTG TCC CAT CAG AGG CTC ATG GAT      2928
Glu Asn Phe Lys Ile Ala Phe Tyr Leu Ser His Gln Arg Leu Met Asp
    940                 945                 950

GAA AAA GTT GAT GTG ACT ATC TTG ACA TTT CTT GAG AAG CCC TTT CAG      2976
Glu Lys Val Asp Val Ile Ile Leu Ile Phe Leu Glu Lys Pro Phe Gln
955                 960                 965                 970

AAG TCC AAG TTC CTC CAG CTC CGG AAA AGG CTC TGT GGG AGT TCT GTC      3024
Lys Ser Lys Phe Leu Gln Leu Arg Lys Arg Leu Cys Gly Ser Ser Val
                975                 980                 985

CTT GAG TGG CCA ACA AAC CCG CAA GCT CAC CCA TAC TTC TGG CAG TGT      3072
Leu Glu Trp Pro Thr Asn Pro Gln Ala His Pro Tyr Phe Trp Gln Cys
            990                 995                 1000

CTA AAG AAC GCC CTG GCC ACA GAC AAT CAT GTG GCC TAT AGT CAG GTG      3120
Leu Lys Asn Ala Leu Ala Thr Asp Asn His Val Ala Tyr Ser Gln Val
        1005                1010                1015

TTC AAG GAA ACG GTC TAG                                              3138
Phe Lys Glu Thr Val
    1020

MWTLKRLILILFNIILISKLLGARWFPKTLPCDVTLDVGKNHVIVDCTDKHLTEIPGGIPTNTTNLTLTINHIP
DISPASFHRLDHLVEIDFRCNCVPIPLGSKNNMCIKRLQIKPRSFSGLTYLKSLYLDGNQLLEIPQGLPPSLQL
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LSLEANNIFSIRKENLTELANIEILYLGQNCYYRNPCYVSYSIEKDAFLNLTKLKVLSLKDNNVTAVPTVLPST
LTELYLYNNMIAKIQEDDFNNLNQLQILDLSGNCPRCYNAPFPCAPCKNNSPLQIPVNAFDALTELKVLRLHSN
SLQHVPPRWFKNINKLQELDLSQNFLAKEIGDAKFLHFLPSLIQLDLSFNFELQVYRASMNLSQAFSSLKSLKI
LRIRGYVFKELKSFNLSPLHNLQNLEVLDLGTNFIKIANLSMFKQFKRLKVIDLSVNKISPSGDSSEVGFCSNA
RTSVESYEPQVLEQLHYFRYDKYARSCRFKNKEASFMSVNESCYKYGQTLDLSKNSIFFVKSSDFQHLSFLKCL
NLSGNLISQTLNGSEFQPLAELRYLDFSNNRLDLLHSTAFEELHKLEVLDISSNSHYFQSEGITHMLNFTKNLK
VLQKLMMNDNDISSSTSRTMESESLRTLEFRGNHLDVLWREGDNRYLQLFKNLLKLEELDISKNSLSFLPSGVF
DGMPPNLKNLSLAKNGLKSFSWKKLQCLKNLETLDLSHNQLTTVPERLSNCSRSLKNLILKNNQIRSLTKYFLQ
DAFQLRYLDLSSNKIQMIQKTSFPENVLNNLKMLLLHHNRFLCTCDAVWFVWWVNHTEVTIPYLATDVTCVGPG
AHKGQSVISLDLYTCELDLTNLILFSLSISVSLFLMVMMTASHLYFWDVWYIYHFCKAKIKGYQRLISPDCCYD
AFIVYDTKDPAVTEWVLAELVAKLEDPREKHFNLCLEERDWLPGQPVLENLSQSIQLSKKTVFVMTDKYAKTEN
FKIAFYLSHQRLMDEKVDVIILIFLEKPFQKSKFLQLRKRLCGSSVLEWPTNPQAHPYFWQCLKNALATDNHVA
YSQVFKETV

rodent (SEQ ID NO: 13 and 14):

```
CTT GGA AAA CCT CTT CAG AAG TCT AAG TTT CTT CAG CTC AGG AAG AGA        48
Leu Gly Lys Pro Leu Gln Lys Ser Lys Phe Leu Gln Leu Arg Lys Arg
 1               5                  10                  15

CTC TGC AGG AGC TCT GTC CTT GAG TGG CCT GCA AAT CCA CAG GCT CAC        96
Leu Cys Arg Ser Ser Val Leu Glu Trp Pro Ala Asn Pro Gln Ala His
            20                  25                  30

CCA TAC TTC TGG CAG TGC CTG AAA AAT GCC CTG ACC ACA GAC AAT CAT       144
Pro Tyr Phe Trp Gln Cys Leu Lys Asn Ala Leu Thr Thr Asp Asn His
        35                  40                  45

GTG GCT TAT AGT CAA ATG TTC AAG GAA ACA GTC TAG                       180
Val Ala Tyr Ser Gln Met Phe Lys Glu Thr Val
    50                  55
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LGKPLQKSKFLQLRKRLCRSSVLEWPANPQAHPYFWQCLKALTTDNHVAYSQMFKECV

additional rodent, e.g., mouse sequences:

upstream (SEQ ID NO: 27 and 28); nucleotides 186, 196, 217, 276, and 300 designated C, each may be A, C, G, or T:

```
TCC TAT TCT ATG GAA AAA GAT GCT TTC CTA TTT ATG AGA AAT TTG AAG        48
Ser Tyr Ser Met Glu Lys Asp Ala Phe Leu Phe Met Arg Asn Leu Lys
 1               5                  10                  15

GTT CTC TCA CTA AAA GAT AAC AAT GTC ACA GCT GTC CCC ACC ACT TTG        96
Val Leu Ser Leu Lys Asp Asn Asn Val Thr Ala Val Pro Thr Thr Leu
            20                  25                  30

CCA CCT AAT TTA CTA GAG CTC TAT CTT TAT AAC AAT ATC ATT AAG AAA       144
Pro Pro Asn Leu Leu Glu Leu Tyr Leu Tyr Asn Asn Ile Ile Lys Lys
        35                  40                  45

ATC CAA GAA AAT GAT TTC AAT AAC CTC AAT GAG TTG CAA GTC CTT GAC       192
Ile Gln Glu Asn Asp Phe Asn Asn Leu Asn Glu Leu Gln Val Leu Asp
    50                  55                  60

CTA CGT GGA AAT TGC CCT CGA TGT CAT AAT GTC CCA TAT CCG TGT ACA       240
Leu Arg Gly Asn Cys Pro Arg Cys His Asn Val Pro Tyr Pro Cys Thr
65                  70                  75                  80

CCG TGT GAA AAT AAT TCC CCC TTA CAG ATC CAT GAC AAT GCT TTC AAT       288
Pro Cys Glu Asn Asn Ser Pro Leu Gln Ile His Asp Asn Ala Phe Asn
                85                  90                  95

TCA TCG ACA GAC                                                      300
Ser Ser Thr Asp
            100
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SYSMEKDAFLFMRNLKVLSLKDNNVTAVPTTLPPNLLELYLYNNIIKKIQENDFNNLNELQXLDLXGNCPRCXNV
PYPCTPCENNSPLQIHXNAFNSSTX

downstream (SEQ ID NO: 29 and 30); nucleotide 1643 designated A, may be A or G; nucleotide 1664 designated C, may be A, C, G, or T; nucleotides 1690 and 1735 designated G, may be G or T; nucleotide 1719 designated C, may be C or T; and nucleotide 1727 designated A, may be A, G, or T:

```
TCT CCA GAA ATT CCC TGG AAT TCC TTG CCT CCT GAG GTT TTT GAG GGT          48
Ser Pro Glu Ile Pro Trp Asn Ser Leu Pro Pro Glu Val Phe Glu Gly
  1               5                  10                  15

ATG CCG CCA AAT CTA AAG AAT CTC TCC TTG GCC AAA AAT GGG CTC AAA          96
Met Pro Pro Asn Leu Lys Asn Leu Ser Leu Ala Lys Asn Gly Leu Lys
            20                  25                  30

TCT TTC TTT TGG GAC AGA CTC CAG TTA CTG AAG CAT TTG GAA ATT TTG         144
Ser Phe Phe Trp Asp Arg Leu Gln Leu Leu Lys His Leu Glu Ile Leu
        35                  40                  45

GAC CTC AGC CAT AAC CAG CTG ACA AAA GTA CCT GAG AGA TTG GCC AAC         192
Asp Leu Ser His Asn Gln Leu Thr Lys Val Pro Glu Arg Leu Ala Asn
    50                  55                  60

TGT TCC AAA AGT CTC ACA ACA CTG ATT CTT AAG CAT AAT CAA ATC AGG         240
Cys Ser Lys Ser Leu Thr Thr Leu Ile Leu Lys His Asn Gln Ile Arg
65                  70                  75                  80

CAA TTG ACA AAA TAT TTT CTA GAA GAT GCT TTG CAA TTG CGC TAT CTA         288
Gln Leu Thr Lys Tyr Phe Leu Glu Asp Ala Leu Gln Leu Arg Tyr Leu
                85                  90                  95

GAC ATC AGT TCA AAT AAA ATC CAG GTC ATT CAG AAG ACT AGC TTC CCA         336
Asp Ile Ser Ser Asn Lys Ile Gln Val Ile Gln Lys Thr Ser Phe Pro
            100                 105                 110

GAA AAT GTC CTC AAC AAT CTG GAG ATG TTG GTT TTA CAT CAC AAT CGC         384
Glu Asn Val Leu Asn Asn Leu Glu Met Leu Val Leu His His Asn Arg
        115                 120                 125

TTT CTT TGC AAC TGT GAT GCT GTG TGG TTT GTC TGG TGG GTT AAC CAT         432
Phe Leu Cys Asn Cys Asp Ala Val Trp Phe Val Trp Trp Val Asn His
    130                 135                 140

ACA GAT GTT ACT ATT CCA TAC CTG GCC ACT GAT GTG ACT TGT GTA GGT         480
Thr Asp Val Thr Ile Pro Tyr Leu Ala Thr Asp Val Thr Cys Val Gly
145                 150                 155                 160

CCA GGA GCA CAC AAA GGT CAA AGT GTC ATA TCC CTT GAT CTG TAT ACG         528
Pro Gly Ala His Lys Gly Gln Ser Val Ile Ser Leu Asp Leu Tyr Thr
                165                 170                 175
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TGT GAG TTA GAT CTC ACA AAC CTG ATT CTG TTC TCA GTT TCC ATA TCA      576
Cys Glu Leu Asp Leu Thr Asn Leu Ile Leu Phe Ser Val Ser Ile Ser
            180                 185                 190

TCA GTC CTC TTT CTT ATG GTA GTT ATG ACA ACA AGT CAC CTC TTT TTC      624
Ser Val Leu Phe Leu Met Val Val Met Thr Thr Ser His Leu Phe Phe
        195                 200                 205

TGG GAT ATG TGG TAC ATT TAT TAT TTT TGG AAA GCA AAG ATA AAG GGG      672
Trp Asp Met Trp Tyr Ile Tyr Tyr Phe Trp Lys Ala Lys Ile Lys Gly
    210                 215                 220

TAT CCA GCA TCT GCA ATC CCA TGG AGT CCT TGT TAT GAT GCT TTT ATT      720
Tyr Pro Ala Ser Ala Ile Pro Trp Ser Pro Cys Tyr Asp Ala Phe Ile
225                 230                 235                 240

GTG TAT GAC ACT AAA AAC TCA GCT GTG ACA GAA TGG GTT TTG CAG GAG      768
Val Tyr Asp Thr Lys Asn Ser Ala Val Thr Glu Trp Val Leu Gln Glu
                245                 250                 255

CTG GTG GCA AAA TTG GAA GAT CCA AGA GAA AAA CAC TTC AAT TTG TGT      816
Leu Val Ala Lys Leu Glu Asp Pro Arg Glu Lys His Phe Asn Leu Cys
            260                 265                 270

CTA GAA GAA AGA GAC TGG CTA CCA GGA CAG CCA GTT CTA GAA AAC CTT      864
Leu Glu Glu Arg Asp Trp Leu Pro Gly Gln Pro Val Leu Glu Asn Leu
        275                 280                 285

TCC CAG AGC ATA CAG CTC AGC AAA AAG ACA GTG TTT GTG ATG ACA CAG      912
Ser Gln Ser Ile Gln Leu Ser Lys Lys Thr Val Phe Val Met Thr Gln
    290                 295                 300

AAA TAT GCT AAG ACT GAG AGT TTT AAG ATG GCA TTT TAT TTG TCT CAT      960
Lys Tyr Ala Lys Thr Glu Ser Phe Lys Met Ala Phe Tyr Leu Ser His
305                 310                 315                 320

CAG AGG CTC CTG GAT GAA AAA GTG GAT GTG ATT ATC TTG ATA TTC TTG     1008
Gln Arg Leu Leu Asp Glu Lys Val Asp Val Ile Ile Leu Ile Phe Leu
                325                 330                 335

GAA AGA CCT CTT CAG AAG TCT AAG TTT CTT CAG CTC AGG AAG AGA CTC     1056
Glu Arg Pro Leu Gln Lys Ser Lys Phe Leu Gln Leu Arg Lys Arg Leu
            340                 345                 350

TGC AGG AGC TCT GTC CTT GAG TGG CCT GCA AAT CCA CAG GCT CAC CCA     1104
Cys Arg Ser Ser Val Leu Glu Trp Pro Ala Asn Pro Gln Ala His Pro
        355                 360                 365

TAC TTC TGG CAG TGC CTG AAA AAT GCC CTG ACC ACA GAC AAT CAT GTG     1152
Tyr Phe Trp Gln Cys Leu Lys Asn Ala Leu Thr Thr Asp Asn His Val
    370                 375                 380

GCT TAT AGT CAA ATG TTC AAG GAA ACA GTC TAGCTCTCTG AAGAATGTCA      1202
Ala Tyr Ser Gln Met Phe Lys Glu Thr Val
385                 390
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CCACCTAGGA CATGCCTTGG TACCTGAAGT TTTCATAAAG GTTTCCATAA ATGAAGGTCT 1262

GAATTTTTCC TAACAGTTGT CATGGCTCAG ATTGGTGGGA AATCATCAAT ATATGGCTAA 1322

GAAATTAAGA AGGGGAGACT GATAGAAGAT AATTTCTTTC TTCATGTGCC ATGCTCAGTT 1382

AAATATTTCC CCTAGCTCAA ATCTGAAAAA CTGTGCCTAG GAGACAACAC AAGGCTTTGA 1442

TTTATCTGCA TACAATTGAT AAGAGCCACA CATCTGCCCT GAAGAAGTAC TAGTAGTTTT 1502

AGTAGTAGGG TAAAAATTAC ACAAGCTTTC TCTCTCTCTG ATACTGAACT GTACCAGAGT 1562

TCAATGAAAT AAAAGCCCAG AGAACTTCTC AGTAAATGGT TTCATTATCA TGTAGTATCC 1622

ACCATGCAAT ATGCCACAAA ACCGCTACTG GTACAGGACA GCTGGTAGCT GCTTCAAGGC 1682

CTCTTATCAT TTTCTTGGGG CCCATGGAGC GGTTCTCTGG GAAAAAGGGA AGGTTTTTTT 1742

TGGCCATCCA TGAA 1756

SPEIPWNSLPPEVFEGMPPNLKNLSLAKNGLKSFFWDRLQLLKHLEILDLSHNQLTKVPERLANCSKSLTTLILK
HNQIRQLTKYFLEDALQLRYLDISSNKIQVIQKTSFPENVLNNLEMLVLHHNRFLCNCDAVWFVWWVNHTDVTIP
YLATDVTCVGPGAHKGQSVISLDLYTCELDLTNLILFSVSISSVLFLMVVMTTSHLFFWDMWYIYYFWKAKIKGY
PASAIPWSPCYDAFIVYDTKNSAVTEWVLQELVAKLEDPREKHFNLCLEERDWLPGQPVLENLSQSIQLSKKTVF
VMTQKYAKTESFKMAFYLSHQRLLDEKVDVIILIFLERPLQKSKFLQLRKRLCRSSVLEWPANPQAHPYFWQCLK
NALTTDNHVAYSQMFKETV

Table 7: Nucleotide and amino acid sequences of a mammalian, e.g., primate, human, DNAX Toll like Receptor 7 (DTLR7).

upstream (SEQ ID NO: 15 and 16):

```
G AAT TCC AGA CTT ATA AAC TTG AAA AAT CTC TAT TTG GCC TGG AAC          46
  Asn Ser Arg Leu Ile Asn Leu Lys Asn Leu Tyr Leu Ala Trp Asn
    1               5                  10                  15

TGC TAT TTT AAC AAA GTT TGC GAG AAA ACT AAC ATA GAA GAT GGA GTA         94
Cys Tyr Phe Asn Lys Val Cys Glu Lys Thr Asn Ile Glu Asp Gly Val
                20                  25                  30

TTT GAA ACG CTG ACA AAT TTG GAG TTG CTA TCA CTA TCT TTC AAT TCT        142
Phe Glu Thr Leu Thr Asn Leu Glu Leu Leu Ser Leu Ser Phe Asn Ser
            35                  40                  45

CTT TCA CAT GTG CCA CCC AAA CTG CCA AGC TCC CTA CGC AAA CTT TTT        190
Leu Ser His Val Pro Pro Lys Leu Pro Ser Ser Leu Arg Lys Leu Phe
        50                  55                  60

CTG AGC AAC ACC CAG ATC AAA TAC ATT AGT GAA GAA GAT TTC AAG GGA        238
Leu Ser Asn Thr Gln Ile Lys Tyr Ile Ser Glu Glu Asp Phe Lys Gly
    65                  70                  75

TTG ATA AAT TTA ACA TTA CTA GAT TTA AGC GGG AAC TGT CCG AGG TGC        286
Leu Ile Asn Leu Thr Leu Leu Asp Leu Ser Gly Asn Cys Pro Arg Cys
80                  85                  90                  95

TTC AAT GCC CCA TTT CCA TGC GTG CCT TGT GAT GGT GGT GCT TCA ATT        334
Phe Asn Ala Pro Phe Pro Cys Val Pro Cys Asp Gly Gly Ala Ser Ile
                100                 105                 110

AAT ATA GAT CGT TTT GCT TTT CAA AAC TTG ACC CAA CTT CGA TAC CTA        382
Asn Ile Asp Arg Phe Ala Phe Gln Asn Leu Thr Gln Leu Arg Tyr Leu
            115                 120                 125

AAC CTC TCT AGC ACT TCC CTC AGG AAG ATT AAT GCT GCC TGG TTT AAA        430
Asn Leu Ser Ser Thr Ser Leu Arg Lys Ile Asn Ala Ala Trp Phe Lys
        130                 135                 140

AAT ATG CCT CAT CTG AAG GTG CTG GAT CTT GAA TTC AAC TAT TTA GTG        478
Asn Met Pro His Leu Lys Val Leu Asp Leu Glu Phe Asn Tyr Leu Val
    145                 150                 155

GGA GAA ATA GCC TCT GGG GCA TTT TTA ACG ATG CTG CCC CGC TTA GAA        526
Gly Glu Ile Ala Ser Gly Ala Phe Leu Thr Met Leu Pro Arg Leu Glu
160                 165                 170                 175

ATA CTT GAC TTG TCT TTT AAC TAT ATA AAG GGG AGT TAT CCA CAG CAT        574
Ile Leu Asp Leu Ser Phe Asn Tyr Ile Lys Gly Ser Tyr Pro Gln His
                180                 185                 190
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ATT AAT ATT TCC AGA AAC TTC TCT AAA CTT TTG TCT CTA CGG GCA TTG      622
Ile Asn Ile Ser Arg Asn Phe Ser Lys Leu Leu Ser Leu Arg Ala Leu
            195                 200                 205

CAT TTA AGA GGT TAT GTG TTC CAG GAA CTC AGA GAA GAT GAT TTC CAG      670
His Leu Arg Gly Tyr Val Phe Gln Glu Leu Arg Glu Asp Asp Phe Gln
        210                 215                 220

CCC CTG ATG CAG CTT CCA AAC TTA TCG ACT ATC AAC TTG GGT ATT AAT      718
Pro Leu Met Gln Leu Pro Asn Leu Ser Thr Ile Asn Leu Gly Ile Asn
    225                 230                 235

TTT ATT AAG CAA ATC GAT TTC AAA CTT TTC CAA AAT TTC TCC AAT CTG      766
Phe Ile Lys Gln Ile Asp Phe Lys Leu Phe Gln Asn Phe Ser Asn Leu
240                 245                 250                 255

GAA ATT ATT TAC TTG TCA GAA AAC AGA ATA TCA CCG TTG GTA AAA GAT      814
Glu Ile Ile Tyr Leu Ser Glu Asn Arg Ile Ser Pro Leu Val Lys Asp
                260                 265                 270

ACC CGG CAG AGT TAT GCA AAT AGT TCC TCT TTT CAA CGT CAT ATC CGG      862
Thr Arg Gln Ser Tyr Ala Asn Ser Ser Ser Phe Gln Arg His Ile Arg
            275                 280                 285

AAA CGA CGC TCA ACA GAT TTT GAG TTT GAC CCA CAT TCG AAC TTT TAT      910
Lys Arg Arg Ser Thr Asp Phe Glu Phe Asp Pro His Ser Asn Phe Tyr
        290                 295                 300

CAT TTC ACC CGT CCT TTA ATA AAG CCA CAA TGT GCT GCT TAT GGA AAA      958
His Phe Thr Arg Pro Leu Ile Lys Pro Gln Cys Ala Ala Tyr Gly Lys
    305                 310                 315

GCC TTA GAT TTA AGC CTC AAC AGT ATT TTC TT                          990
Ala Leu Asp Leu Ser Leu Asn Ser Ile Phe
320                 325

NSRLINLKNLYLAWNCYFNKVCEKTNIEDGVFETLTNLELLSLSFNSLSHVPPKLPSSLRKLFLSNTQIKYISE
EDFKGLINLTLLDLSGNCPRCFNAPFPCVPCDGGASINIDRFAFQNLTQLRYLNLSSTSLRKINAAWFKNMPHL
KVLDLEFNYLVGEIASGAFLTMLPRLEILDLSFNYIKGSYPQHINISRNFSKLLSLRALHLRGYVFQELREDDF
QPLMQLPNLSTINLGINFIKQIDFKLFQNFSNLEIIYLSENRISPLVKDTRQSYANSSSFQRHIRKRRSTDFEF
DPHSNFYHFTRPLIKPQCAAYGKALDLSLNSIF
```

downstream (SEQ ID NO: 17 and 18):

```
CAG TCT CTT TCC ACA TCC CAA ACT TTC TAT GAT GCT TAC ATT TCT TAT       48
Gln Ser Leu Ser Thr Ser Gln Thr Phe Tyr Asp Ala Tyr Ile Ser Tyr
 1               5                  10                  15

GAC ACC AAA GAT GCC TCT GTT ACT GAC TGG GTG ATA AAT GAG CTG CGC       96
Asp Thr Lys Asp Ala Ser Val Thr Asp Trp Val Ile Asn Glu Leu Arg
            20                  25                  30
```

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TAC CAC CTT GAA GAG AGC CGA GAC AAA AAC GTT CTC CTT TGT CTA GAG      144
Tyr His Leu Glu Glu Ser Arg Asp Lys Asn Val Leu Leu Cys Leu Glu
         35                  40                  45

GAG AGG GAT TGG GAC CCG GGA TTG GCC ATC ATC GAC AAC CTC ATG CAG      192
Glu Arg Asp Trp Asp Pro Gly Leu Ala Ile Ile Asp Asn Leu Met Gln
     50                  55                  60

AGC ATC AAC CAA AGC AAG AAA ACA GTA TTT GTT TTA ACC AAA AAA TAT      240
Ser Ile Asn Gln Ser Lys Lys Thr Val Phe Val Leu Thr Lys Lys Tyr
 65                  70                  75                  80

GCA AAA AGC TGG AAC TTT AAA ACA GCT TTT TAC TTG GGC TTG CAG AGG      288
Ala Lys Ser Trp Asn Phe Lys Thr Ala Phe Tyr Leu Gly Leu Gln Arg
                 85                  90                  95

CTA ATG GGT GAG AAC ATG GAT GTG ATT ATA TTT ATC CTG CTG GAG CCA      336
Leu Met Gly Glu Asn Met Asp Val Ile Ile Phe Ile Leu Leu Glu Pro
            100                 105                 110

GTG TTA CAG CAT TCT CCG TAT TTG AGG CTA CGG CAG CGG ATC TGT AAG      384
Val Leu Gln His Ser Pro Tyr Leu Arg Leu Arg Gln Arg Ile Cys Lys
        115                 120                 125

AGC TCC ATC CTC CAG TGG CCT GAC AAC CCG AAG GCA GAA AGG TTG TTT      432
Ser Ser Ile Leu Gln Trp Pro Asp Asn Pro Lys Ala Glu Arg Leu Phe
    130                 135                 140

TGG CAA ACT CTG AGA AAT GTG GTC TTG ACT GAA AAT GAT TCA CGG TAT      480
Trp Gln Thr Leu Arg Asn Val Val Leu Thr Glu Asn Asp Ser Arg Tyr
145                 150                 155                 160

AAC AAT ATG TAT GTC GAT TCC ATT AAG CAA TAC TAACTGACGT TAAGTCATGA     533
Asn Asn Met Tyr Val Asp Ser Ile Lys Gln Tyr
                165                 170

TTTCGCGCCA TAATAAAGAT GCAAAGGAAT GACATTTCCG TATTAGTTAT CTATTGCTAC     593

GGTAACCAAA TTACTCCCAA AAACCTTACG TCGGTTTCAA AACAACCACA TTCTGCTGGC     653

CCCACAGTTT TTGAGGGTCA GGAGTCCAGG CCCAGCATAA CTGGGTCTTC TGCTTCAGGG     713

TGTCTCCAGA GGCTGCAATG TAGGTGTTCA CCAGAGACAT AGGCATCACT GGGGTCACAC     773

TCCATGTGGT TGTTTTCTGG ATTCAATTCC TCCTGGGCTA TTGGCCAAAG GCTATACTCA     833

TGTAAGCCAT GCGAGCCTAT CCCACAACGG CAGCTTGCTT CATCAGAGCT AGCAAAAAAG     893

AGAGGTTGCT AGCAAGATGA AGTCACAATC TTTTGTAATC GAATCAAAAA AGTGATATCT     953

CATCACTTTG GCCATATTCT ATTTGTTAGA AGTAAACCAC AGGTCCCACC AGCTCCATGG    1013

GAGTGACCAC CTCAGTCCAG GGAAAACAGC TGAAGACCAA GATGGTGAGC TCTGATTGCT    1073

TCAGTTGGTC ATCAACTATT TTCCCTTGAC TGCTGTCCTG GGATGGCCGG CTATCTTGAT    1133
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GGATAGATTG TGAATATCAG GAGGCCAGGG ATCACTGTGG ACCATCTTAG CAGTTGACCT    1193

AACACATCTT CTTTTCAATA TCTAAGAACT TTTGCCACTG TGACTAATGG TCCTAATATT    1253

AAGCTGTTGT TTATATTTAT CATATATCTA TGGCTACATG GTTATATTAT GCTGTGGTTG    1313

CGTTCGGTTT TATTTACAGT TGCTTTTACA AATATTTGCT GTAACATTTG ACTTCTAAGG    1373

TTTAGATGCC ATTTAAGAAC TGAGATGGAT AGCTTTTAAA GCATCTTTTA CTTCTTACCA    1433

TTTTTTAAAA GTATGCAGCT AAATTCGAAG CTTTTGGTCT ATATTGTTAA TTGCCATTGC    1493

TGTAAATCTT AAAATGAATG AATAAAAATG TTTCATTTTA AAAAAAAAAA AAAAAAAAAA    1553

AAAA                                                                 1557
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QSLSTSQTFYDAYISYDTKDASVTDWVINELRYHLEESRDKNVLLCLEERDWDPGLAIIDNLMQSINQSKKTVFV
LTKKYAKSWNFKTAFYLGLQRLMGENMDVIIFILLEPVLQHSPYLRLRQRICKSSILQWPDNPKAERLFWQTLRN
VVLTENDSRYNNMYVDSIKQY
```

Further primate, e.g, human, DTLR7 sequence (SEQ ID NO: 36 and 37).

```
atg ctg acc tgc att ttc ctg cta ata tct ggt tcc tgt gag tta tgc   48
Met Leu Thr Cys Ile Phe Leu Leu Ile Ser Gly Ser Cys Glu Leu Cys
        -15                 -10                 -5

gcc gaa gaa aat ttt tct aga agc tat cct tgt gat gag aaa aag caa   96
Ala Glu Glu Asn Phe Ser Arg Ser Tyr Pro Cys Asp Glu Lys Lys Gln
 -1   1               5                  10                  15

aat gac tca gtt att gca gag tgc agc aat cgt cga cta cag gaa gtt   144
Asn Asp Ser Val Ile Ala Glu Cys Ser Asn Arg Arg Leu Gln Glu Val
                20                  25                  30

ccc caa acg gtg ggc aaa tat gtg aca gaa cta gac ctg tct gat aat   192
Pro Gln Thr Val Gly Lys Tyr Val Thr Glu Leu Asp Leu Ser Asp Asn
            35                  40                  45

ttc atc aca cac ata acg aat gaa tca ttt caa ggg ctg caa aat ctc   240
Phe Ile Thr His Ile Thr Asn Glu Ser Phe Gln Gly Leu Gln Asn Leu
        50                  55                  60

act aaa ata aat cta aac cac aac ccc aat gta cag cac cag aac gga   288
Thr Lys Ile Asn Leu Asn His Asn Pro Asn Val Gln His Gln Asn Gly
    65                  70                  75

aat ccc ggt ata caa tca aat ggc ttg aat atc aca gac ggg gca ttc   336
Asn Pro Gly Ile Gln Ser Asn Gly Leu Asn Ile Thr Asp Gly Ala Phe
80                  85                  90                  95

ctc aac cta aaa aac cta agg gag tta ctg ctt gaa gac aac cag tta   384
Leu Asn Leu Lys Asn Leu Arg Glu Leu Leu Leu Glu Asp Asn Gln Leu
                100                 105                 110
```

```
ccc caa ata ccc tct ggt ttg cca gag tct ttg aca gaa ctt agt cta   432
Pro Gln Ile Pro Ser Gly Leu Pro Glu Ser Leu Thr Glu Leu Ser Leu
            115                 120                 125

att caa aac aat ata tac aac ata act aaa gag ggc att tca aga ctt   480
Ile Gln Asn Asn Ile Tyr Asn Ile Thr Lys Glu Gly Ile Ser Arg Leu
        130                 135                 140

ata aac ttg aaa aat ctc tat ttg gcc tgg aac tgc tat ttt aac aaa   528
Ile Asn Leu Lys Asn Leu Tyr Leu Ala Trp Asn Cys Tyr Phe Asn Lys
    145                 150                 155

gtt tgc gag aaa act aac ata gaa gat gga gta ttt gaa acg ctg aca   576
Val Cys Glu Lys Thr Asn Ile Glu Asp Gly Val Phe Glu Thr Leu Thr
160                 165                 170                 175

aat ttg gag ttg cta tca cta tct ttc aat tct ctt tca cat gtg cca   624
Asn Leu Glu Leu Leu Ser Leu Ser Phe Asn Ser Leu Ser His Val Pro
                180                 185                 190

ccc aaa ctg cca agc tcc cta cgc aaa ctt ttt ctg agc aac acc cag   672
Pro Lys Leu Pro Ser Ser Leu Arg Lys Leu Phe Leu Ser Asn Thr Gln
            195                 200                 205

atc aaa tac att agt gaa gaa gat ttc aag gga ttg ata aat tta aca   720
Ile Lys Tyr Ile Ser Glu Glu Asp Phe Lys Gly Leu Ile Asn Leu Thr
        210                 215                 220

tta cta gat tta agc ggg aac tgt ccg agg tgc ttc aat gcc cca ttt   768
Leu Leu Asp Leu Ser Gly Asn Cys Pro Arg Cys Phe Asn Ala Pro Phe
    225                 230                 235

cca tgc gtg cct tgt gat ggt ggt gct tca att aat ata gat cgt ttt   816
Pro Cys Val Pro Cys Asp Gly Gly Ala Ser Ile Asn Ile Asp Arg Phe
240                 245                 250                 255

gct ttt caa aac ttg acc caa ctt cga tac cta aac ctc tct agc act   864
Ala Phe Gln Asn Leu Thr Gln Leu Arg Tyr Leu Asn Leu Ser Ser Thr
                260                 265                 270

tcc ctc agg aag att aat gct gcc tgg ttt aaa aat atg cct cat ctg   912
Ser Leu Arg Lys Ile Asn Ala Ala Trp Phe Lys Asn Met Pro His Leu
            275                 280                 285

aag gtg ctg gat ctt gaa ttc aac tat tta gtg gga gaa ata gcc tct   960
Lys Val Leu Asp Leu Glu Phe Asn Tyr Leu Val Gly Glu Ile Ala Ser
        290                 295                 300

ggg gca ttt tta acg atg ctg ccc cgc tta gaa ata ctt gac ttg tct  1008
Gly Ala Phe Leu Thr Met Leu Pro Arg Leu Glu Ile Leu Asp Leu Ser
    305                 310                 315

ttt aac tat ata aag ggg agt tat cca cag cat att aat att tcc aga  1056
Phe Asn Tyr Ile Lys Gly Ser Tyr Pro Gln His Ile Asn Ile Ser Arg
320                 325                 330                 335
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aac ttc tct aaa ctt ttg tct cta cgg gca ttg cat tta aga ggt tat   1104
Asn Phe Ser Lys Leu Leu Ser Leu Arg Ala Leu His Leu Arg Gly Tyr
                340                 345                 350

gtg ttc cag gaa ctc aga gaa gat gat ttc cag ccc ctg atg cag ctt   1152
Val Phe Gln Glu Leu Arg Glu Asp Asp Phe Gln Pro Leu Met Gln Leu
            355                 360                 365

cca aac tta tcg act atc aac ttg ggt att aat ttt att aag caa atc   1200
Pro Asn Leu Ser Thr Ile Asn Leu Gly Ile Asn Phe Ile Lys Gln Ile
        370                 375                 380

gat ttc aaa ctt ttc caa aat ttc tcc aat ctg gaa att att tac ttg   1248
Asp Phe Lys Leu Phe Gln Asn Phe Ser Asn Leu Glu Ile Ile Tyr Leu
    385                 390                 395

tca gaa aac aga ata tca ccg ttg gta aaa gat acc cgg cag agt tat   1296
Ser Glu Asn Arg Ile Ser Pro Leu Val Lys Asp Thr Arg Gln Ser Tyr
400                 405                 410                 415

gca aat agt tcc tct ttt caa cgt cat atc cgg aaa cga cgc tca aca   1344
Ala Asn Ser Ser Ser Phe Gln Arg His Ile Arg Lys Arg Arg Ser Thr
                420                 425                 430

gat ttt gag ttt gac cca cat tcg aac ttt tat cat ttc acc cgt cct   1392
Asp Phe Glu Phe Asp Pro His Ser Asn Phe Tyr His Phe Thr Arg Pro
            435                 440                 445

tta ata aag cca caa tgt gct gct tat gga aaa gcc tta gat tta agc   1440
Leu Ile Lys Pro Gln Cys Ala Ala Tyr Gly Lys Ala Leu Asp Leu Ser
        450                 455                 460

ctc aac agt att ttc ttc att ggg cca aac caa ttt gaa aat ctt cct   1488
Leu Asn Ser Ile Phe Phe Ile Gly Pro Asn Gln Phe Glu Asn Leu Pro
    465                 470                 475

gac att gcc tgt tta aat ctg tct gca aat agc aat gct caa gtg tta   1536
Asp Ile Ala Cys Leu Asn Leu Ser Ala Asn Ser Asn Ala Gln Val Leu
480                 485                 490                 495

agt gga act gaa ttt tca gcc att cct cat gtc aaa tat ttg gat ttg   1584
Ser Gly Thr Glu Phe Ser Ala Ile Pro His Val Lys Tyr Leu Asp Leu
                500                 505                 510

aca aac aat aga cta gac ttt gat aat gct agt gct ctt act gaa ttg   1632
Thr Asn Asn Arg Leu Asp Phe Asp Asn Ala Ser Ala Leu Thr Glu Leu
            515                 520                 525

tcc gac ttg gaa gtt cta gat ctc agc tat aat tca cac tat ttc aga   1680
Ser Asp Leu Glu Val Leu Asp Leu Ser Tyr Asn Ser His Tyr Phe Arg
        530                 535                 540

ata gca ggc gta aca cat cat cta gaa ttt att caa aat ttc aca aat   1728
Ile Ala Gly Val Thr His His Leu Glu Phe Ile Gln Asn Phe Thr Asn
    545                 550                 555

cta aaa gtt tta aac ttg agc cac aac aac att tat act tta aca gat   1776
Leu Lys Val Leu Asn Leu Ser His Asn Asn Ile Tyr Thr Leu Thr Asp
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     560                 565                 570                 575

     aag tat aac ctg gaa agc aag tcc ctg gta gaa tta gtt ttc agt ggc   1824
     Lys Tyr Asn Leu Glu Ser Lys Ser Leu Val Glu Leu Val Phe Ser Gly
 5                   580                 585                 590

     aat cgc ctt gac att ttg tgg aat gat gat gac aac agg tat atc tcc   1872
     Asn Arg Leu Asp Ile Leu Trp Asn Asp Asp Asp Asn Arg Tyr Ile Ser
                 595                 600                 605
10
```

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att ttc aaa ggt ctc aag aat ctg aca cgt ctg gat tta tcc ctt aat   1920
Ile Phe Lys Gly Leu Lys Asn Leu Thr Arg Leu Asp Leu Ser Leu Asn
        610                 615                 620

agg ctc aag cac atc cca aat gaa gca ttc ctt aat ttg cca gcg agt   1968
Arg Leu Lys His Ile Pro Asn Glu Ala Phe Leu Asn Leu Pro Ala Ser
    625                 630                 635

ctc act gaa cta cat ata aat gat aat atg tta aag ttt ttt aac tgg   2016
Leu Thr Glu Leu His Ile Asn Asp Asn Met Leu Lys Phe Phe Asn Trp
640                 645                 650                 655

aca tta ctc cag cag ttt cct cgt ctc gag ttg ctt gac tta cgt gga   2064
Thr Leu Leu Gln Gln Phe Pro Arg Leu Glu Leu Leu Asp Leu Arg Gly
                660                 665                 670

aac aaa cta ctc ttt tta act gat agc cta tct gac ttt aca tct tcc   2112
Asn Lys Leu Leu Phe Leu Thr Asp Ser Leu Ser Asp Phe Thr Ser Ser
            675                 680                 685

ctt cgg aca ctg ctg ctg agt cat aac agg att tcc cac cta ccc tct   2160
Leu Arg Thr Leu Leu Leu Ser His Asn Arg Ile Ser His Leu Pro Ser
        690                 695                 700

ggc ttt ctt tct gaa gtc agt agt ctg aag cac ctc gat tta agt tcc   2208
Gly Phe Leu Ser Glu Val Ser Ser Leu Lys His Leu Asp Leu Ser Ser
    705                 710                 715

aat ctg cta aaa aca atm aac aaa tcc gca ctt gaa act aag acc acc   2256
Asn Leu Leu Lys Thr Xaa Asn Lys Ser Ala Leu Glu Thr Lys Thr Thr
720                 725                 730                 735

acc aaa tta tct atg ttg gaa cta cac gga aac ccc ttt gaa tgc acc   2304
Thr Lys Leu Ser Met Leu Glu Leu His Gly Asn Pro Phe Glu Cys Thr
                740                 745                 750

tgt gac att gga gat ttc cga aga tgg atg gat gaa cat ctg aat gtc   2352
Cys Asp Ile Gly Asp Phe Arg Arg Trp Met Asp Glu His Leu Asn Val
            755                 760                 765

aaa att ccc aga ctg gta gat gtc att tgt gcc agt cct ggg gat caa   2400
Lys Ile Pro Arg Leu Val Asp Val Ile Cys Ala Ser Pro Gly Asp Gln
        770                 775                 780

aga ggg aag agt att gtg agt ctg gag cta aca act tgt gtt tca gat   2448
Arg Gly Lys Ser Ile Val Ser Leu Glu Leu Thr Thr Cys Val Ser Asp
    785                 790                 795

gtc act gca gtg ata tta ttt ttc ttc acg ttc ttt atc acc acc atg   2496
Val Thr Ala Val Ile Leu Phe Phe Phe Thr Phe Phe Ile Thr Thr Met
800                 805                 810                 815

gtt atg ttg gct gcc ctg gct cac cat ttg ttt tac tgg gat gtt tgg   2544
Val Met Leu Ala Ala Leu Ala His His Leu Phe Tyr Trp Asp Val Trp
                820                 825                 830
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ttt ata tat aat gtg tgt tta gct aag tta aaa ggc tac agg tct ctt   2592
Phe Ile Tyr Asn Val Cys Leu Ala Lys Leu Lys Gly Tyr Arg Ser Leu
            835                 840                 845

tcc aca tcc caa act ttc tat gat gct tac att tct tat gac acc aaa   2640
Ser Thr Ser Gln Thr Phe Tyr Asp Ala Tyr Ile Ser Tyr Asp Thr Lys
        850                 855                 860

gat gcc tct gtt act gac tgg gtg ata aat gag ctg cgc tac cac ctt   2688
Asp Ala Ser Val Thr Asp Trp Val Ile Asn Glu Leu Arg Tyr His Leu
    865                 870                 875

gaa gag agc cga gac aaa aac gtt ctc ctt tgt cta gag gag agg gat   2736
Glu Glu Ser Arg Asp Lys Asn Val Leu Leu Cys Leu Glu Glu Arg Asp
880                 885                 890                 895

tgg gac ccg gga ttg gcc atc atc gac aac ctc atg cag agc atc aac   2784
Trp Asp Pro Gly Leu Ala Ile Ile Asp Asn Leu Met Gln Ser Ile Asn
                900                 905                 910

caa agc aag aaa aca gta ttt gtt tta acc aaa aaa tat gca aaa agc   2832
Gln Ser Lys Lys Thr Val Phe Val Leu Thr Lys Lys Tyr Ala Lys Ser
            915                 920                 925

tgg aac ttt aaa aca gct ttt tac ttg gcc ttg cag agg cta atg ggt   2880
Trp Asn Phe Lys Thr Ala Phe Tyr Leu Ala Leu Gln Arg Leu Met Gly
        930                 935                 940

gag aac atg gat gtg att ata ttt atc ctg ctg gag cca gtg tta cag   2928
Glu Asn Met Asp Val Ile Ile Phe Ile Leu Leu Glu Pro Val Leu Gln
    945                 950                 955

cat tct ccg tat ttg agg cta cgg cag cgg atc tgt aag agc tcc atc   2976
His Ser Pro Tyr Leu Arg Leu Arg Gln Arg Ile Cys Lys Ser Ser Ile
960                 965                 970                 975

ctc cag tgg cct gac aac ccg aag gca gaa ggc ttg ttt tgg caa act   3024
Leu Gln Trp Pro Asp Asn Pro Lys Ala Glu Gly Leu Phe Trp Gln Thr
                980                 985                 990

ctg aga aat gtg gtc ttg act gaa aat gat tca cgg tat aac aat atg   3072
Leu Arg Asn Val Val Leu Thr Glu Asn Asp Ser Arg Tyr Asn Asn Met
            995                 1000                1005

tat gtc gat tcc att aag caa tac taa                              3099
Tyr Val Asp Ser Ile Lys Gln Tyr
        1010                1015
```

MLTCIFLLISGSCELCAEENFSRSYPCDEKKQNDSVIAECSNRRLQEVPQTVGKYVTELDLSDNFITHI
TNESFQGLQNLTKINLNHNPNVQHQNGNPGIQSNGLNITDGAFLNLKNLRELLLEDNQLPQIPSGLPES
LTELSLIQNNIYNITKEGISRLINLKNLYLAWNCYFNKVCEKTNIEDGVFETLTNLELLSLSFNSLSHV
PPKLPSSLRKLFLSNTQIKYISEEDFKGLINLTLLDLSGNCPRCFNAPFPCVPCDGGASINIDRFAFQN
LTQLRYLNLSSTSLRKINAAWFKNMPHLKVLDLEFNYLVGEIASGAFLTMLPRLEILDLSFNYIKGSYP
QHINISRNFSKLLSLRALHLRGYVFQELREDDFQPLMQLPNLSTINLGINFIKQIDFKLFQNFSNLEII
YLSENRISPLVKDTRQSYANSSSFQRHIRKRRSTDFEFDPHSNFYHFTRPLIKPQCAAYGKALDLSLNS
IFFIGPNQFENLPDIACLNLSANSNAQVLSGTEFSAIPHVKYLDLTNNRLDFDNASALTELSDLEVLDL
SYNSHYFRIAGVTHHLEFIQNFTNLKVLNLSHNNIYTLTDKYNLESKSLVELVFSGNRLDILWNDDDNR
YISIFKGLKNLTRLDLSLNRLKHIPNEAFLNLPASLTELHINDNMLKFFNWTLLQQFPRLELLDLRGNK
LLFLTDSLSDFTSSLRTLLLSHNRISHLPSGFLSEVSSLKHLDLSSNLLKTINKSALETKTTTKLSMLE
LHGNPFECTCDIGDFRRWMDEHLNVKIPRLVDVICASPGDQRGKSIVSLELTTCVSDVTAVILFFFTFF
ITTMVMLAALAHHLFYWDVWFIYNVCLAKLKGYRSLSTSQTFYDAYISYDTKDASVTDWVINELRYHLE
ESRDKNVLLCLEERDWDPGLAIIDNLMQSINQSKKTVFVLTKKYAKSWNFKTAFYLALQRLMGENMDVI
IFILLEPVLQHSPYLRLRQRICKSSILQWPDNPKAEGLFWQTLRNVVLTENDSRYNNMYVDSIKQY

Table 8: Partial nucleotide and amino acid sequences (see SEQ ID NO: 19 and 20) of a mammalian, e.g., primate, human, DNAX Toll like Receptor 8 (DTLR8).

```
AAT GAA TTG ATC CCC AAT CTA GAG AAG GAA GAT GGT TCT ATC TTG ATT          48
Asn Glu Leu Ile Pro Asn Leu Glu Lys Glu Asp Gly Ser Ile Leu Ile
 1               5                  10                  15

TGC CTT TAT GAA AGC TAC TTT GAC CCT GGC AAA AGC ATT AGT GAA AAT          96
Cys Leu Tyr Glu Ser Tyr Phe Asp Pro Gly Lys Ser Ile Ser Glu Asn
            20                  25                  30

ATT GTA AGC TTC ATT GAG AAA AGC TAT AAG TCC ATC TTT GTT TTG TCC         144
Ile Val Ser Phe Ile Glu Lys Ser Tyr Lys Ser Ile Phe Val Leu Ser
        35                  40                  45

CCC AAC TTT GTC CAG AAT GAG TGG TGC CAT TAT GAA TTC TAC TTT GCC         192
Pro Asn Phe Val Gln Asn Glu Trp Cys His Tyr Glu Phe Tyr Phe Ala
    50                  55                  60

CAC CAC AAT CTC TTC CAT GAA AAT TCT GAT CAC ATA ATT CTT ATC TTA         240
His His Asn Leu Phe His Glu Asn Ser Asp His Ile Ile Leu Ile Leu
65                  70                  75                  80

CTG GAA CCC ATT CCA TTC TAT TGC ATT CCC ACC AGG TAT CAT AAA CTG         288
Leu Glu Pro Ile Pro Phe Tyr Cys Ile Pro Thr Arg Tyr His Lys Leu
                85                  90                  95

GAA GCT CTC CTG GAA AAA AAA GCA TAC TTG GAA TGG CCC AAG GAT AGG         336
Glu Ala Leu Leu Glu Lys Lys Ala Tyr Leu Glu Trp Pro Lys Asp Arg
            100                 105                 110

CGT AAA TGT GGG CTT TTC TGG GCA AAC CTT CGA GCT GCT GTT AAT GTT         384
Arg Lys Cys Gly Leu Phe Trp Ala Asn Leu Arg Ala Ala Val Asn Val
        115                 120                 125

AAT GTA TTA GCC ACC AGA GAA ATG TAT GAA CTG CAG ACA TTC ACA GAG         432
Asn Val Leu Ala Thr Arg Glu Met Tyr Glu Leu Gln Thr Phe Thr Glu
    130                 135                 140

TTA AAT GAA GAG TCT CGA GGT TCT ACA ATC TCT CTG ATG AGA ACA GAC         480
Leu Asn Glu Glu Ser Arg Gly Ser Thr Ile Ser Leu Met Arg Thr Asp
145                 150                 155                 160

TGT CTA TAAAATCCCA CAGTCCTTGG GAAGTTGGGG ACCACATACA CTGTTGGGAT          536
Cys Leu

GTACATTGAT ACAACCTTTA TGATGGCAAT TTGACAATAT TTATTAAAAT AAAAAATGGT       596

TATTCCCTTC AAAAAAAAAA AAAAAAAAAA AAA                                    629

NELIPNLEKEDGSILICLYESYFDEGKSISENIVSFIEKSYKSIFVLSPNFVQNEWCHYEFYFAHHNLFHENS
HIILILLEPIPFYCIPTRYHKLEALLEKKAYLEWPKDRRKCGLFWANLRAAVNVNVLATREMYELQTFTELNI
SRGSTISLMRTDCL
```

additional primate, e.g., human sequence (SEQ ID NO: 31 and 32); nucleotides 4 and 23 designated C, may be A, C, G, or T; nucleotide 845 designated C, may be C or T:

```
C TCC GAT GCC AAG ATT CGG CAC CAG GCA TAT TCA GAG GTC ATG ATG          46
  Ser Asp Ala Lys Ile Arg His Gln Ala Tyr Ser Glu Val Met Met
    1               5                  10                  15

GTT GGA TGG TCA GAT TCA TAC ACC TGT GAA TAC CCT TTA AAC CTA AGG        94
Val Gly Trp Ser Asp Ser Tyr Thr Cys Glu Tyr Pro Leu Asn Leu Arg
                  20                  25                  30

GGA ACT AGG TTA AAA GAC GTT CAT CTC CAC GAA TTA TCT TGC AAC ACA       142
Gly Thr Arg Leu Lys Asp Val His Leu His Glu Leu Ser Cys Asn Thr
              35                  40                  45

GCT CTG TTG ATT GTC ACC ATT GTG GTT ATT ATG CTA GTT CTG GGG TTG       190
Ala Leu Leu Ile Val Thr Ile Val Val Ile Met Leu Val Leu Gly Leu
          50                  55                  60

GCT GTG GCC TTC TGC TGT CTC CAC TTT GAT CTG CCC TGG TAT CTC AGG       238
Ala Val Ala Phe Cys Cys Leu His Phe Asp Leu Pro Trp Tyr Leu Arg
      65                  70                  75

ATG CTA GGT CAA TGC ACA CAA ACA TGG CAC AGG GTT AGG AAA ACA ACC       286
Met Leu Gly Gln Cys Thr Gln Thr Trp His Arg Val Arg Lys Thr Thr
  80                  85                  90                  95

CAA GAA CAA CTC AAG AGA AAT GTC CGA TTC CAC GCA TTT ATT TCA TAC       334
Gln Glu Gln Leu Lys Arg Asn Val Arg Phe His Ala Phe Ile Ser Tyr
                 100                 105                 110

AGT GAA CAT GAT TCT CTG TGG GTG AAG AAT GAA TTG ATC CCC AAT CTA       382
Ser Glu His Asp Ser Leu Trp Val Lys Asn Glu Leu Ile Pro Asn Leu
             115                 120                 125

GAG AAG GAA GAT GGT TCT ATC TTG ATT TGC CTT TAT GAA AGC TAC TTT       430
Glu Lys Glu Asp Gly Ser Ile Leu Ile Cys Leu Tyr Glu Ser Tyr Phe
         130                 135                 140

GAC CCT GGC AAA AGC ATT AGT GAA AAT ATT GTA AGC TTC ATT GAG AAA       478
Asp Pro Gly Lys Ser Ile Ser Glu Asn Ile Val Ser Phe Ile Glu Lys
     145                 150                 155

AGC TAT AAG TCC ATC TTT GTT TTG TCT CCC AAC TTT GTC CAG AAT GAG       526
Ser Tyr Lys Ser Ile Phe Val Leu Ser Pro Asn Phe Val Gln Asn Glu
160                 165                 170                 175

TGG TGC CAT TAT GAA TTC TAC TTT GCC CAC CAC AAT CTC TTC CAT GAA       574
Trp Cys His Tyr Glu Phe Tyr Phe Ala His His Asn Leu Phe His Glu
                 180                 185                 190

AAT TCT GAT CAC ATA ATT CTT ATC TTA CTG GAA CCC ATT CCA TTC TAT       622
Asn Ser Asp His Ile Ile Leu Ile Leu Leu Glu Pro Ile Pro Phe Tyr
             195                 200                 205
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TGC ATT CCC ACC AGG TAT CAT AAA CTG GAA GCT CTC CTG GAA AAA AAA      670
Cys Ile Pro Thr Arg Tyr His Lys Leu Glu Ala Leu Leu Glu Lys Lys
        210                 215                 220

GCA TAC TTG GAA TGG CCC AAG GAT AGG CGT AAA TGT GGG CTT TTC TGG      718
Ala Tyr Leu Glu Trp Pro Lys Asp Arg Arg Lys Cys Gly Leu Phe Trp
    225                 230                 235

GCA AAC CTT CGA GCT GCT GTT AAT GTT AAT GTA TTA GCC ACC AGA GAA      766
Ala Asn Leu Arg Ala Ala Val Asn Val Asn Val Leu Ala Thr Arg Glu
240                 245                 250                 255

ATG TAT GAA CTG CAG ACA TTC ACA GAG TTA AAT GAA GAG TCT CGA GGT      814
Met Tyr Glu Leu Gln Thr Phe Thr Glu Leu Asn Glu Glu Ser Arg Gly
                260                 265                 270

TCT ACA ATC TCT CTG ATG AGA ACA GAC TGT CTA TAAAATCCCA CAGTCCTTGG    867
Ser Thr Ile Ser Leu Met Arg Thr Asp Cys Leu
            275                 280

GAAGTTGGGG ACCACATACA CTGTTGGGAT GTACATTGAT ACAACCTTTA TGATGGCAAT    927

TTGACAATAT TTATTAAAAT AAAAAATGGT TATTCCCTTC AAAAAAAAAA AAAAAAAAAA    987

AAAAAAAAAA AA                                                       999

SDAKIRHQAYSEVMMVGWSDSYTCEYPLNLRGTRLKDVHLHELSCNTALLIVTIVVIMLVLGLAVAFCCLHFDI
WYLRMLGQCTQTWHRVRKTTQEQLKRNVRFHAFISYSEHDSLWVKNELIPNLEKEDGSILICLYESYFDPGKSI
ENIVSFIEKSYKSIFVLSPNFVQNEWCHYEFYFAHHNLFHENSDHIILILLEPIPFYCIPTRYHKLEALLEKKA
LEWPKDRRKCGLFWANLRAAVKVNVLATREMYELQTFTELNEESRGSTISLMRTDCL
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Further primate, e.g., human, DTLR8 (SEQ ID NO: 38 and 39):

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gaatcatcca cgcacctgca gctctgctga gagagtgcaa gccgtggggg ttttgagctc 60

atcttcatca ttcatatgag gaaataagtg gtaaaatcct tggaaataca atg aga     116
                                                        Met Arg

ctc atc aga aac att tac ata ttt tgt agt att gtt atg aca gca gag    164
Leu Ile Arg Asn Ile Tyr Ile Phe Cys Ser Ile Val Met Thr Ala Glu
        -15                 -10                 -5

ggt gat gct cca gag ctg cca gaa gaa agg gaa ctg atg acc aac tgc    212
Gly Asp Ala Pro Glu Leu Pro Glu Glu Arg Glu Leu Met Thr Asn Cys
 -1   1               5                  10                  15

tcc aac atg tct cta aga aag gtt ccc gca gac ttg acc cca gcc aca    260
Ser Asn Met Ser Leu Arg Lys Val Pro Ala Asp Leu Thr Pro Ala Thr
                20                  25                  30

acg aca ctg gat tta tcc tat aac ctc ctt ttt caa ctc cag agt tca    308
Thr Thr Leu Asp Leu Ser Tyr Asn Leu Leu Phe Gln Leu Gln Ser Ser
            35                  40                  45
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gat ttt cat tct gtc tcc aaa ctg aga gtt ttg att cta tgc cat aac   356
Asp Phe His Ser Val Ser Lys Leu Arg Val Leu Ile Leu Cys His Asn
          50                  55                  60

aga att caa cag ctg gat ctc aaa acc ttt gaa ttc aac aag gag tta   404
Arg Ile Gln Gln Leu Asp Leu Lys Thr Phe Glu Phe Asn Lys Glu Leu
      65                  70                  75

aga tat tta gat ttg tct aat aac aga ctg aag agt gta act tgg tat   452
Arg Tyr Leu Asp Leu Ser Asn Asn Arg Leu Lys Ser Val Thr Trp Tyr
  80                  85                  90                  95

tta ctg gca ggt ctc agg tat tta gat ctt tct ttt aat gac ttt gac   500
Leu Leu Ala Gly Leu Arg Tyr Leu Asp Leu Ser Phe Asn Asp Phe Asp
              100                 105                 110

acc atg cct atc tgt gag gaa gct ggc aac atg tca cac ctg gaa atc   548
Thr Met Pro Ile Cys Glu Glu Ala Gly Asn Met Ser His Leu Glu Ile
          115                 120                 125

cta ggt ttg agt ggg gca aaa ata caa aaa tca gat ttc cag aaa att   596
Leu Gly Leu Ser Gly Ala Lys Ile Gln Lys Ser Asp Phe Gln Lys Ile
      130                 135                 140

gct cat ctg cat cta aat act gtc ttc tta gga ttc aga act ctt cct   644
Ala His Leu His Leu Asn Thr Val Phe Leu Gly Phe Arg Thr Leu Pro
  145                 150                 155

cat tat gaa gaa ggt agc ctg ccc atc tta aac aca aca aaa ctg cac   692
His Tyr Glu Glu Gly Ser Leu Pro Ile Leu Asn Thr Thr Lys Leu His
160                 165                 170                 175

att gtt tta cca atg gac aca aat ttc tgg gtt ctt ttg cgt gat gga   740
Ile Val Leu Pro Met Asp Thr Asn Phe Trp Val Leu Leu Arg Asp Gly
              180                 185                 190

atc aag act tca aaa ata tta gaa atg aca aat ata gat ggc aaa agc   788
Ile Lys Thr Ser Lys Ile Leu Glu Met Thr Asn Ile Asp Gly Lys Ser
          195                 200                 205

caa ttt gta agt tat gaa atg caa cga aat ctt agt tta gaa aat gct   836
Gln Phe Val Ser Tyr Glu Met Gln Arg Asn Leu Ser Leu Glu Asn Ala
      210                 215                 220

aag aca tcg gtt cta ttg ctt aat aaa gtt gat tta ctc tgg gac gac   884
Lys Thr Ser Val Leu Leu Leu Asn Lys Val Asp Leu Leu Trp Asp Asp
  225                 230                 235

ctt ttc ctt atc tta caa ttt gtt tgg cat aca tca gtg gaa cac ttt   932
Leu Phe Leu Ile Leu Gln Phe Val Trp His Thr Ser Val Glu His Phe
240                 245                 250                 255

cag atc cga aat gtg act ttt ggt ggt aag gct tat ctt gac cac aat   980
Gln Ile Arg Asn Val Thr Phe Gly Gly Lys Ala Tyr Leu Asp His Asn
              260                 265                 270
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tca ttt gac tac tca aat act gta atg aga act ata aaa ttg gag cat   1028
Ser Phe Asp Tyr Ser Asn Thr Val Met Arg Thr Ile Lys Leu Glu His
            275                 280                 285

gta cat ttc aga gtg ttt tac att caa cag gat aaa atc tat ttg ctt   1076
Val His Phe Arg Val Phe Tyr Ile Gln Gln Asp Lys Ile Tyr Leu Leu
        290                 295                 300

ttg acc aaa atg gac ata gaa aac ctg aca ata tca aat gca caa atg   1124
Leu Thr Lys Met Asp Ile Glu Asn Leu Thr Ile Ser Asn Ala Gln Met
    305                 310                 315

cca cac atg ctt ttc ccg aat tat cct acg aaa ttc caa tat tta aat   1172
Pro His Met Leu Phe Pro Asn Tyr Pro Thr Lys Phe Gln Tyr Leu Asn
320                 325                 330                 335

ttt gcc aat aat atc tta aca gac gag ttg ttt aaa aga act atc caa   1220
Phe Ala Asn Asn Ile Leu Thr Asp Glu Leu Phe Lys Arg Thr Ile Gln
                340                 345                 350

ctg cct cac ttg aaa act ctc att ttg aat ggc aat aaa ctg gag aca   1268
Leu Pro His Leu Lys Thr Leu Ile Leu Asn Gly Asn Lys Leu Glu Thr
            355                 360                 365

ctt tct tta gta agt tgc ttt gct aac aac aca ccc ttg gaa cac ttg   1316
Leu Ser Leu Val Ser Cys Phe Ala Asn Asn Thr Pro Leu Glu His Leu
        370                 375                 380

gat ctg agt caa aat cta tta caa cat aaa aat gat gaa aat tgc tca   1364
Asp Leu Ser Gln Asn Leu Leu Gln His Lys Asn Asp Glu Asn Cys Ser
    385                 390                 395

tgg cca gaa act gtg gtc aat atg aat ctg tca tac aat aaa ttg tct   1412
Trp Pro Glu Thr Val Val Asn Met Asn Leu Ser Tyr Asn Lys Leu Ser
400                 405                 410                 415

gat tct gtc ttc agg tgc ttg ccc aaa agt att caa ata ctt gac cta   1460
Asp Ser Val Phe Arg Cys Leu Pro Lys Ser Ile Gln Ile Leu Asp Leu
                420                 425                 430

aat aat aac caa atc caa act gta cct aaa gag act att cat ctg atg   1508
Asn Asn Asn Gln Ile Gln Thr Val Pro Lys Glu Thr Ile His Leu Met
            435                 440                 445

gcc tta cga gaa cta aat att gca ttt aat ttt cta act gat ctc cct   1556
Ala Leu Arg Glu Leu Asn Ile Ala Phe Asn Phe Leu Thr Asp Leu Pro
        450                 455                 460

gga tgc agt cat ttc agt aga ctt tca gtt ctg aac att gaa atg aac   1604
Gly Cys Ser His Phe Ser Arg Leu Ser Val Leu Asn Ile Glu Met Asn
    465                 470                 475

ttc att ctc agc cca tct ctg gat ttt gtt cag agc tgc cag gaa gtt   1652
Phe Ile Leu Ser Pro Ser Leu Asp Phe Val Gln Ser Cys Gln Glu Val
480                 485                 490                 495
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aaa act cta aat gcg gga aga aat cca ttc cgg tgt acc tgt gaa tta 1700
Lys Thr Leu Asn Ala Gly Arg Asn Pro Phe Arg Cys Thr Cys Glu Leu
                500                 505                 510

aaa aat ttc att cag ctt gaa aca tat tca gag gtc atg atg gtt gga 1748
Lys Asn Phe Ile Gln Leu Glu Thr Tyr Ser Glu Val Met Met Val Gly
            515                 520                 525

tgg tca gat tca tac acc tgt gaa tac cct tta aac cta agg gga act 1796
Trp Ser Asp Ser Tyr Thr Cys Glu Tyr Pro Leu Asn Leu Arg Gly Thr
        530                 535                 540

agg tta aaa gac gtt cat ctc cac gaa tta tct tgc aac aca gct ctg 1844
Arg Leu Lys Asp Val His Leu His Glu Leu Ser Cys Asn Thr Ala Leu
    545                 550                 555

ttg att gtc acc att gtg gtt att atg cta gtt ctg ggg ttg gct gtg 1892
Leu Ile Val Thr Ile Val Val Ile Met Leu Val Leu Gly Leu Ala Val
560                 565                 570                 575

gcc ttc tgc tgt ctc cac ttt gat ctg ccc tgg tat ctc agg atg cta 1940
Ala Phe Cys Cys Leu His Phe Asp Leu Pro Trp Tyr Leu Arg Met Leu
                580                 585                 590

ggt caa tgc aca caa aca tgg cac agg gtt agg aaa aca acc caa gaa 1988
Gly Gln Cys Thr Gln Thr Trp His Arg Val Arg Lys Thr Thr Gln Glu
            595                 600                 605

caa ctc aag aga aat gtc cga ttc cac gca ttt att tca tac agt gaa 2036
Gln Leu Lys Arg Asn Val Arg Phe His Ala Phe Ile Ser Tyr Ser Glu
        610                 615                 620

cat gat tct ctg tgg gtg aag aat gaa ttg atc ccc aat cta gag aag 2084
His Asp Ser Leu Trp Val Lys Asn Glu Leu Ile Pro Asn Leu Glu Lys
    625                 630                 635

gaa gat ggt tct atc ttg att tgc ctt tat gaa agc tac ttt gac cct 2132
Glu Asp Gly Ser Ile Leu Ile Cys Leu Tyr Glu Ser Tyr Phe Asp Pro
640                 645                 650                 655

ggc aaa agc att agt gaa aat att gta agc ttc att gag aaa agc tat 2180
Gly Lys Ser Ile Ser Glu Asn Ile Val Ser Phe Ile Glu Lys Ser Tyr
                660                 665                 670

aag tcc atc ttt gtt ttg tct ccc aac ttt gtc cag aat gag tgg tgc 2228
Lys Ser Ile Phe Val Leu Ser Pro Asn Phe Val Gln Asn Glu Trp Cys
            675                 680                 685

cat tat gaa ttc tac ttt gcc cac cac aat ctc ttc cat gaa aat tct 2276
His Tyr Glu Phe Tyr Phe Ala His His Asn Leu Phe His Glu Asn Ser
        690                 695                 700

gat cat ata att ctt atc tta ctg gaa ccc att cca ttc tat tgc att 2324
Asp His Ile Ile Leu Ile Leu Leu Glu Pro Ile Pro Phe Tyr Cys Ile
    705                 710                 715
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ccc acc agg tat cat aaa ctg aaa gct ctc ctg gaa aaa aaa gca tac 2372
Pro Thr Arg Tyr His Lys Leu Lys Ala Leu Leu Glu Lys Lys Ala Tyr
720 725 730 735

ttg gaa tgg ccc aag gat agg cgt aaa tgt ggg ctt ttc tgg gca aac 2420
Leu Glu Trp Pro Lys Asp Arg Arg Lys Cys Gly Leu Phe Trp Ala Asn
740 745 750

ctt cga gct gct att aat gtt aat gta tta gcc acc aga gaa atg tat 2468
Leu Arg Ala Ala Ile Asn Val Asn Val Leu Ala Thr Arg Glu Met Tyr
755 760 765

gaa ctg cag aca ttc aca gag tta aat gaa gag tct cga ggt tct aca 2516
Glu Leu Gln Thr Phe Thr Glu Leu Asn Glu Glu Ser Arg Gly Ser Thr
770 775 780

atc tct ctg atg aga aca gat tgt cta taaaatccca cagtccttgg 2563
Ile Ser Leu Met Arg Thr Asp Cys Leu
785 790

gaagttgggg accacataca ctgttgggat gtacattgat acaaccttta tgatggcaat 2623

ttgacaatat ttattaaaat aaaaaatggt tattcccttc atatcagttt ctagaaggat 2683

ttctaagaat gtatcctata gaaacacctt cacaagttta taagggctta tggaaaaagg 2743

tgttcatccc aggattgttt ataatcatga aaaatgtggc caggtgcagt ggctcactct 2803

tgtaatccca gcactatggg aggccaaggt gggtgaccca cgaggtcaag agatggagac 2863

catcctggcc aacatggtga aaccctgtct ctactaaaaa tacaaaaatt agctgggcgt 2923

gatggtgcac gcctgtagtc ccagctactt gggaggctga ggcaggagaa tcgcttgaac 2983

ccgggaggtg gcagttgcag tgagctgaga tcgagccact gcactccagc ctggtgacag 3043

agc 3046

MRLIRNIYIFCSIVMTAEGDAPELPEERELMTNCSNMSLRKVPADLTPATTTLDLSYNLLFQLQSSDFH
SVSKLRVLILCHNRIQQLDLKTFEFNKELRYLDLSNNRLKSVTWYLLAGLRYLDLSFNDFDTMPICEEA
GNMSHLEILGLSGAKIQKSDFQKIAHLHLNTVFLGFRTLPHYEEGSLPILNTTKLHIVLPMDTNFWVLL
RDGIKTSKILEMTNIDGKSQFVSYEMQRNLSLENAKTSVLLLNKVDLLWDDLFLILQFVWHTSVEHFQI
RNVTFGGKAYLDHNSFDYSNTVMRTIKLEHVHFRVFYIQQDKIYLLLTKMDIENLTISNAQMPHMLFPN
YPTKFQYLNFANNILTDELFKRTIQLPHLKTLILNGNKLETLSLVSCFANNTPLEHLDLSQNLLQHKND
ENCSWPETVVNMNLSYNKLSDSVFRCLPKSIQILDLNNNQIQTVPKETIHLMALRELNIAFNFLTDLPG
CSHFSRLSVLNIEMNFILSPSLDFVQSCQEVKTLNAGRNPFRCTCELKNFIQLETYSEVMMVGWSDSYT
CEYPLNLRGTRLKDVHLHELSCNTALLIVTIVVIMLVLGLAVAFCCLHFDLPWYLRMLGQCTQTWHRVR
KTTQEQLKRNVRFHAFISYSEHDSLWVKNELIPNLEKEDGSILICLYESYFDPGKSISENIVSFIEKSY
KSIFVLSPNFVQNEWCHYEFYFAHHNLFHENSDHIILILLEPIPFYCIPTRYHKLKALLEKKAYLEWPK
DRRKCGLFWANLRAAINVNVLATREMYELQTFTELNEESRGSTISLMRTDCL

Table 9: Partial nucleotide and amino acid sequences (see SEQ ID NO: 21 and 22) of a mammalian, e.g., primate, human, DNAX Toll like Receptor 9 (DTLR9).

```
AAG AAC TCC AAA GAA AAC CTC CAG TTT CAT GCT TTT ATT TCA TAT AGT          48
Lys Asn Ser Lys Glu Asn Leu Gln Phe His Ala Phe Ile Ser Tyr Ser
 1               5                  10                  15

GAA CAT GAT TCT GCC TGG GTG AAA AGT GAA TTG GTA CCT TAC CTA GAA          96
Glu His Asp Ser Ala Trp Val Lys Ser Glu Leu Val Pro Tyr Leu Glu
            20                  25                  30

AAA GAA GAT ATA CAG ATT TGT CTT CAT GAG AGA AAC TTT GTC CCT GGC         144
Lys Glu Asp Ile Gln Ile Cys Leu His Glu Arg Asn Phe Val Pro Gly
        35                  40                  45

AAG AGC ATT GTG GAA AAT ATC ATC AAC TGC ATT GAG AAG AGT TAC AAG         192
Lys Ser Ile Val Glu Asn Ile Ile Asn Cys Ile Glu Lys Ser Tyr Lys
    50                  55                  60

TCC ATC TTT GTT TTG TCT CCC AAC TTT GTC CAG AGT GAG TGG TGC CAT         240
Ser Ile Phe Val Leu Ser Pro Asn Phe Val Gln Ser Glu Trp Cys His
65                  70                  75                  80

TAC GAA CTC TAT TTT GCC CAT CAC AAT CTC TTT CAT GAA GGA TCT AAT         288
Tyr Glu Leu Tyr Phe Ala His His Asn Leu Phe His Glu Gly Ser Asn
                85                  90                  95

AAC TTA ATC CTC ATC TTA CTG GAA CCC ATT CCA CAG AAC AGC ATT CCC         336
Asn Leu Ile Leu Ile Leu Leu Glu Pro Ile Pro Gln Asn Ser Ile Pro
            100                 105                 110

AAC AAG TAC CAC AAG CTG AAG GCT CTC ATG ACG CAG CGG ACT TAT TTG         384
Asn Lys Tyr His Lys Leu Lys Ala Leu Met Thr Gln Arg Thr Tyr Leu
        115                 120                 125

CAG TGG CCC AAG GAG AAA AGC AAA CGT GGG CTC TTT TGG GCT                 426
Gln Trp Pro Lys Glu Lys Ser Lys Arg Gly Leu Phe Trp Ala
    130                 135                 140

A                                                                       427

KNSKENLQFHAFISYSEHDSAWVKSELVPYLEKEDIQICLHERNFVPGKSIVENIINCIEKSYKSIFVLSPNF
SEWCHYELYFAHHNLFHEGSNNLILILLEPIPQNSIPNKYHKLKALMTQRTYLQWPKEKSKRGLFWA
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Further primate, e.g., human DTLR9 (SEQ ID NO: 40 and 41):

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aagaatttgg actcatatca agatgctctg aagaagaaca acccttttagg atagccactg 60

caacatc atg acc aaa gac aaa gaa cct att gtt aaa agc ttc cat ttt 109
       Met Thr Lys Asp Lys Glu Pro Ile Val Lys Ser Phe His Phe
          -30                 -25                 -20

gtt tgc ctt atg atc ata ata gtt gga acc aga atc cag ttc tcc gac 157
Val Cys Leu Met Ile Ile Ile Val Gly Thr Arg Ile Gln Phe Ser Asp
       -15                 -10                 -5

gga aat gaa ttt gca gta gac aag tca aaa aga ggt ctt att cat gtt 205
Gly Asn Glu Phe Ala Val Asp Lys Ser Lys Arg Gly Leu Ile His Val
 -1  1                5                  10                  15

cca aaa gac cta ccg ctg aaa acc aaa gtc tta gat atg tct cag aac 253
Pro Lys Asp Leu Pro Leu Lys Thr Lys Val Leu Asp Met Ser Gln Asn
                20                  25                  30

tac atc gct gag ctt cag gtc tct gac atg agc ttt cta tca gag ttg 301
Tyr Ile Ala Glu Leu Gln Val Ser Asp Met Ser Phe Leu Ser Glu Leu
            35                  40                  45

aca gtt ttg aga ctt tcc cat aac aga atc cag cta ctt gat tta agt 349
Thr Val Leu Arg Leu Ser His Asn Arg Ile Gln Leu Leu Asp Leu Ser
        50                  55                  60

gtt ttc aag ttc aac cag gat tta gaa tat ttg gat tta tct cat aat 397
Val Phe Lys Phe Asn Gln Asp Leu Glu Tyr Leu Asp Leu Ser His Asn
    65                  70                  75

cag ttg caa aag ata tcc tgc cat cct att gtg agt ttc agg cat tta 445
Gln Leu Gln Lys Ile Ser Cys His Pro Ile Val Ser Phe Arg His Leu
80                  85                  90                  95

gat ctc tca ttc aat gat ttc aag gcc ctg ccc atc tgt aag gaa ttt 493
Asp Leu Ser Phe Asn Asp Phe Lys Ala Leu Pro Ile Cys Lys Glu Phe
                100                 105                 110

ggc aac tta tca caa ctg aat ttc ttg gga ttg agt gct atg aag ctg 541
Gly Asn Leu Ser Gln Leu Asn Phe Leu Gly Leu Ser Ala Met Lys Leu
            115                 120                 125

caa aaa tta gat ttg ctg cca att gct cac ttg cat cta agt tat atc 589
Gln Lys Leu Asp Leu Leu Pro Ile Ala His Leu His Leu Ser Tyr Ile
        130                 135                 140

ctt ctg gat tta aga aat tat tat ata aaa gaa aat gag aca gaa agt 637
Leu Leu Asp Leu Arg Asn Tyr Tyr Ile Lys Glu Asn Glu Thr Glu Ser
    145                 150                 155

cta caa att ctg aat gca aaa acc ctt cac ctt gtt ttt cac cca act 685
Leu Gln Ile Leu Asn Ala Lys Thr Leu His Leu Val Phe His Pro Thr
160                 165                 170                 175
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agt tta ttc gct atc caa gtg aac ata tca gtt aat act tta ggg tgc   733
Ser Leu Phe Ala Ile Gln Val Asn Ile Ser Val Asn Thr Leu Gly Cys
                180                 185                 190

tta caa ctg act aat att aaa ttg aat gat gac aac tgt caa gtt ttc   781
Leu Gln Leu Thr Asn Ile Lys Leu Asn Asp Asp Asn Cys Gln Val Phe
            195                 200                 205

att aaa ttt tta tca gaa ctc acc aga ggt cca acc tta ctg aat ttt   829
Ile Lys Phe Leu Ser Glu Leu Thr Arg Gly Pro Thr Leu Leu Asn Phe
        210                 215                 220

acc ctc aac cac ata gaa acg act tgg aaa tgc ctg gtc aga gtc ttt   877
Thr Leu Asn His Ile Glu Thr Thr Trp Lys Cys Leu Val Arg Val Phe
    225                 230                 235

caa ttt ctt tgg ccc aaa cct gtg gaa tat ctc aat att tac aat tta   925
Gln Phe Leu Trp Pro Lys Pro Val Glu Tyr Leu Asn Ile Tyr Asn Leu
240                 245                 250                 255

aca ata att gaa agc att cgt gaa gaa gat ttt act tat tct aaa acg   973
Thr Ile Ile Glu Ser Ile Arg Glu Glu Asp Phe Thr Tyr Ser Lys Thr
                260                 265                 270

aca ttg aaa gca ttg aca ata gaa cat atc acg aac caa gtt ttt ctg   1021
Thr Leu Lys Ala Leu Thr Ile Glu His Ile Thr Asn Gln Val Phe Leu
            275                 280                 285

ttt tca cag aca gct ttg tac acc gtg ttt tct gag atg aac att atg   1069
Phe Ser Gln Thr Ala Leu Tyr Thr Val Phe Ser Glu Met Asn Ile Met
        290                 295                 300

atg tta acc att tca gat aca cct ttt ata cac atg ctg tgt cct cat   1117
Met Leu Thr Ile Ser Asp Thr Pro Phe Ile His Met Leu Cys Pro His
    305                 310                 315

gca cca agc aca ttc aag ttt ttg aac ttt acc cag aac gtt ttc aca   1165
Ala Pro Ser Thr Phe Lys Phe Leu Asn Phe Thr Gln Asn Val Phe Thr
320                 325                 330                 335

gat agt att ttt gaa aaa tgt tcc acg tta gtt aaa ttg gag aca ctt   1213
Asp Ser Ile Phe Glu Lys Cys Ser Thr Leu Val Lys Leu Glu Thr Leu
                340                 345                 350

atc tta caa aag aat gga tta aaa gac ctt ttc aaa gta ggt ctc atg   1261
Ile Leu Gln Lys Asn Gly Leu Lys Asp Leu Phe Lys Val Gly Leu Met
            355                 360                 365

acg aag gat atg cct tct ttg gaa ata ctg gat gtt agc tgg aat tct   1309
Thr Lys Asp Met Pro Ser Leu Glu Ile Leu Asp Val Ser Trp Asn Ser
        370                 375                 380

ttg gaa tct ggt aga cat aaa gaa aac tgc act tgg gtt gag agt ata   1357
Leu Glu Ser Gly Arg His Lys Glu Asn Cys Thr Trp Val Glu Ser Ile
    385                 390                 395
```

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gtg gtg tta aat ttg tct tca aat atg ctt act gac tct gtt ttc aga   1405
Val Val Leu Asn Leu Ser Ser Asn Met Leu Thr Asp Ser Val Phe Arg
400                 405                 410                 415

tgt tta cct ccc agg atc aag gta ctt gat ctt cac agc aat aaa ata   1453
Cys Leu Pro Pro Arg Ile Lys Val Leu Asp Leu His Ser Asn Lys Ile
                420                 425                 430

aag agc gtt cct aaa caa gtc gta aaa ctg gaa gct ttg caa gaa ctc   1501
Lys Ser Val Pro Lys Gln Val Val Lys Leu Glu Ala Leu Gln Glu Leu
            435                 440                 445

aat gtt gct ttc aat tct tta act gac ctt cct gga tgt ggc agc ttt   1549
Asn Val Ala Phe Asn Ser Leu Thr Asp Leu Pro Gly Cys Gly Ser Phe
        450                 455                 460

agc agc ctt tct gta ttg atc att gat cac aat tca gtt tcc cac cca   1597
Ser Ser Leu Ser Val Leu Ile Ile Asp His Asn Ser Val Ser His Pro
    465                 470                 475

tcg gct gat ttc ttc cag agc tgc cag aag atg agg tca ata aaa gca   1645
Ser Ala Asp Phe Phe Gln Ser Cys Gln Lys Met Arg Ser Ile Lys Ala
480                 485                 490                 495

ggg gac aat cca ttc caa tgt acc tgt gag cta aga gaa ttt gtc aaa   1693
Gly Asp Asn Pro Phe Gln Cys Thr Cys Glu Leu Arg Glu Phe Val Lys
                500                 505                 510

aat ata gac caa gta tca agt gaa gtg tta gag ggc tgg cct gat tct   1741
Asn Ile Asp Gln Val Ser Ser Glu Val Leu Glu Gly Trp Pro Asp Ser
            515                 520                 525

tat aag tgt gac tac cca gaa agt tat aga gga agc cca cta aag gac   1789
Tyr Lys Cys Asp Tyr Pro Glu Ser Tyr Arg Gly Ser Pro Leu Lys Asp
        530                 535                 540

ttt cac atg tct gaa tta tcc tgc aac ata act ctg ctg atc gtc acc   1837
Phe His Met Ser Glu Leu Ser Cys Asn Ile Thr Leu Leu Ile Val Thr
    545                 550                 555

atc ggt gcc acc atg ctg gtg ttg gct gtg act gtg acc tcc ctc tgc   1885
Ile Gly Ala Thr Met Leu Val Leu Ala Val Thr Val Thr Ser Leu Cys
560                 565                 570                 575

atc tac ttg gat ctg ccc tgg tat ctc agg atg gtg tgc cag tgg acc   1933
Ile Tyr Leu Asp Leu Pro Trp Tyr Leu Arg Met Val Cys Gln Trp Thr
                580                 585                 590

cag act cgg cgc agg gcc agg aac ata ccc tta gaa gaa ctc caa aga   1981
Gln Thr Arg Arg Arg Ala Arg Asn Ile Pro Leu Glu Glu Leu Gln Arg
            595                 600                 605

aac ctc cag ttt cat gct ttt att tca tat agt gaa cat gat tct gcc   2029
Asn Leu Gln Phe His Ala Phe Ile Ser Tyr Ser Glu His Asp Ser Ala
        610                 615                 620
```

tgg gtg aaa agt gaa ttg gta cct tac cta gaa aaa gaa gat ata cag 2077
Trp Val Lys Ser Glu Leu Val Pro Tyr Leu Glu Lys Glu Asp Ile Gln
625 630 635

att tgt ctt cat gag agg aac ttt gtc cct ggc aag agc att gtg gaa 2125
Ile Cys Leu His Glu Arg Asn Phe Val Pro Gly Lys Ser Ile Val Glu
640 645 650 655

aat atc atc aac tgc att gag aag agt tac aag tcc atc ttt gtt ttg 2173
Asn Ile Ile Asn Cys Ile Glu Lys Ser Tyr Lys Ser Ile Phe Val Leu
660 665 670

tct ccc aac ttt gtc cag agt gag tgg tgc cat tac gaa ctc tat ttt 2221
Ser Pro Asn Phe Val Gln Ser Glu Trp Cys His Tyr Glu Leu Tyr Phe
675 680 685

gcc cat cac aat ctc ttt cat gaa gga tct aat aac tta atc ctc atc 2269
Ala His His Asn Leu Phe His Glu Gly Ser Asn Asn Leu Ile Leu Ile
690 695 700

tta ctg gaa ccc att cca cag aac agc att ccc aac aag tac cac aag 2317
Leu Leu Glu Pro Ile Pro Gln Asn Ser Ile Pro Asn Lys Tyr His Lys
705 710 715

ctg aag gct ctc atg acg cag cgg act tat ttg cag tgg ccc aag gag 2365
Leu Lys Ala Leu Met Thr Gln Arg Thr Tyr Leu Gln Trp Pro Lys Glu
720 725 730 735

aaa agc aaa cgt ggg ctc ttt tgg gct aac att aga gcc gct ttt aat 2413
Lys Ser Lys Arg Gly Leu Phe Trp Ala Asn Ile Arg Ala Ala Phe Asn
740 745 750

atg aaa tta aca cta gtc act gaa aac aat gat gtg aaa tct 2455
Met Lys Leu Thr Leu Val Thr Glu Asn Asn Asp Val Lys Ser
755 760 765

taaaaaaatt taggaaattc aacttaagaa accattattt acttggatga tggtgaatag 2515

tacagtcgta agtnactgtc tggaggtgcc tccattatcc tcatgccttc aggaaagact 2575

taacaaaaac aatgtttcat ctggggaact gagctaggcg gtgaggttag cctgccagtt 2635

agagacagcc cagtctcttc tggtttaatc attatgtttc aaattgaaac agtctctttt 2695

gagtaaatgc tcagtttttc agctcctctc cactctgctt tcccaaatgg attctgttgg 2755

tgaag 2760

MTKDKEPIVKSFHFVCLMIIIVGTRIQFSDGNEFAVDKSKRGLIHVPKDLPLKTKVLDMSQNYIAELQV
SDMSFLSELTVLRLSHNRIQLLDLSVFKFNQDLEYLDLSHNQLQKISCHPIVSFRHLDLSFNDFKALPI
CKEFGNLSQLNFLGLSAMKLQKLDLLPIAHLHLSYILLDLRNYYIKENETESLQILNAKTLHLVFHPTS
LFAIQVNISVNTLGCLQLTNIKLNDDNCQVFIKFLSELTRGPTLLNFTLNHIETTWKCLVRVFQFLWPK
PVEYLNIYNLTIIESIRESDFTYSKTTLKALTIEHITNQVFLFSQTALYTVFSEMNIMMLTISDTPFIH
MLCPHAPSTFKFLNFTQNVFTDSIFEKCSTLVKLETLILQKNGLKDLFKVGLMTKDMPSLEILDVSWNS
LESGRHKENCTWVESIVVLNLSSNMLTDSVFRCLPPRIKVLDLHSNKIKSVPKQVVKLEALQELNVAFN
SLTDLPGCGSFSSLSVLIIDHNSVSHPSADFFQSCQKMRSIKAGDNPFQCTCELREFVKNIDQVSSEVL
EGWPDSYKCDYPESYRGSPLKDFHMSELSCNITLLIVTIGATMLVLAVTVTSLCIYLDLPWYLRMVCQW
TQTRRRARNIPLEELQRNLQFHAFISYSEHDSAWVKSELVPYLEKEDIQICLHERNFVPGKSIVENIIN
CIEKSYKSIFVLSPNFVQSEWCHYELYFAHHNLFHEGSNNLILILLEPIPQNSIPNKYHKLKALMTQRT
YLQWPKEKSKRGLFWANIRAAFNMKLTLVTENNDVKS

Table 10: Nucleotide and amino acid sequences (see SEQ ID NO: 23 and 24) of a mammalian, e.g., primate, human, DNAX Toll like Receptor 10 (DTLR1C). Nucleotides 54, 103, and 345 are designated A; each may be A or G; nucleotide 313 designated G, may be G or T; and nucleotides 316, 380, 407, and 408 designated C; each may be A, C, G, or T.

```
GCT TCC ACC TGT GCC TGG CCT GGC TTC CCT GGC GGG GGC GGC AAA GTG        48
Ala Ser Thr Cys Ala Trp Pro Gly Phe Pro Gly Gly Gly Gly Lys Val
  1               5                  10                  15

GGC GAA ATG AGG ATG CCC TGC CCT ACG ATG CCT TCG TGG TCT TCG ACA        96
Gly Glu Met Arg Met Pro Cys Pro Thr Met Pro Ser Trp Ser Ser Thr
            20                  25                  30

AAA CGC AGA GCG CAG TGG CAG ACT GGG TGT ACA ACG AGC TTC GGG GGC       144
Lys Arg Arg Ala Gln Trp Gln Thr Gly Cys Thr Thr Ser Phe Gly Gly
        35                  40                  45

AGC TGG AGG AGT GCC GTG GGC GCT GGG CAC TCC GCC TGT GCC TGG AGG       192
Ser Trp Arg Ser Ala Val Gly Ala Gly His Ser Ala Cys Ala Trp Arg
    50                  55                  60

AAC GCG ACT GGC TGC CTG GCA AAA CCC TCT TTG AGA ACC TGT GGG CCT       240
Asn Ala Thr Gly Cys Leu Ala Lys Pro Ser Leu Arg Thr Cys Gly Pro
65                  70                  75                  80

CGG TCT ATG GCA GCC GCA AGA CGC TGT TTG TGC TGG CCC ACA CGG ACC       288
Arg Ser Met Ala Ala Ala Arg Arg Cys Leu Cys Trp Pro Thr Arg Thr
                85                  90                  95

GGG TCA GTG GTC TCT TGC GCG CCA GTT CTC CTG CTG GCC CAG CAG CGC       336
Gly Ser Val Val Ser Cys Ala Pro Val Leu Leu Leu Ala Gln Gln Arg
            100                 105                 110

CTG CTG GAA GAC CGC AAG GAC GTC GTG GTG CTG GTG ATC CTA ACG CCT       384
Leu Leu Glu Asp Arg Lys Asp Val Val Val Leu Val Ile Leu Thr Pro
        115                 120                 125

GAC GGC CAA GCC TCC CGA CTA CCC GAT GCG CTG ACC AGC GCC TCT GCC       432
Asp Gly Gln Ala Ser Arg Leu Pro Asp Ala Leu Thr Ser Ala Ser Ala
    130                 135                 140

GCC AGA GTG TCC TCC TCT GGC CCC ACC AGC CCA GTG GTC GCG CAG CTT       480
Ala Arg Val Ser Ser Ser Gly Pro Thr Ser Pro Val Val Ala Gln Leu
145                 150                 155                 160

CTG AGG CCA GCA TGC ATG GCC CTG ACC AGG GAC AAC CAC CAC TTC TAT       528
Leu Arg Pro Ala Cys Met Ala Leu Thr Arg Asp Asn His His Phe Tyr
                165                 170                 175

AAC CGG AAC TTC TGC CAG GGA ACC CAC GGC CGA ATA GCC GTG AGC CGG       576
Asn Arg Asn Phe Cys Gln Gly Thr His Gly Arg Ile Ala Val Ser Arg
            180                 185                 190
```

```
AAT CCT GCA CGG TGC CAC CTC CAC ACA CAC CTA ACA TAT GCC TGC CTG      624
Asn Pro Ala Arg Cys His Leu His Thr His Leu Thr Tyr Ala Cys Leu
        195                 200                 205

ATC TGACCAACAC ATGCTCGCCA CCCTCACCAC ACACC                          662
Ile

ASTCAWPGFPGGGGKVGEMRMPCPTMPSWSSTKRRAQWQTGCTTSFGGSWRSAVGAGHSACAWRNATGCLAKPSL
RTCGPRSMAAARRCLCWPTRTGSVVSCAPVLLLAQQRLLEDRKDVVVLVILTFDGQASRLPDALTSASAARVSSS
GPTSPVVAQLLRPACMALTRDNHHFYNRNFCQGTHGRIAVSRNPARCSLHTHLTYACLI
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additional primate, e.g., human DTLR10 sequence (SEQ ID NO: 33 and 34); nucleotide 854 designated A, may be A or T; and nucleotides 1171 and 1172 designated C, each may be A, C, G, or T:

```
CTG CCT GCT GGC ACC CGG CTC CGG AGG CTG GAT GTC AGC TGC AAC AGC      48
Leu Pro Ala Gly Thr Arg Leu Arg Arg Leu Asp Val Ser Cys Asn Ser
  1               5                  10                  15

ATC AGC TTC GTG GCC CCC GGC TTC TTT TCC AAG GCC AAG GAG CTG CGA      96
Ile Ser Phe Val Ala Pro Gly Phe Phe Ser Lys Ala Lys Glu Leu Arg
            20                  25                  30

GAG CTC AAC CTT AGC GCC AAC GCC CTC AAG ACA GTG GAC CAC TCC TGG     144
Glu Leu Asn Leu Ser Ala Asn Ala Leu Lys Thr Val Asp His Ser Trp
        35                  40                  45

TTT GGG CCC CTG GCG AGT GCC CTG CAA ATA CTA GAT GTA AGC GCC AAC     192
Phe Gly Pro Leu Ala Ser Ala Leu Gln Ile Leu Asp Val Ser Ala Asn
    50                  55                  60

CCT CTG CAC TGC GCC TGT GGG GCG GCC TTT ATG GAC TTC CTG CTG GAG     240
Pro Leu His Cys Ala Cys Gly Ala Ala Phe Met Asp Phe Leu Leu Glu
65                  70                  75                  80

GTG CAG GCT GCC GTC CCC GGT CTG CCC AGC CGG GTG AAG TGT GGC AGT     288
Val Gln Ala Ala Val Pro Gly Leu Pro Ser Arg Val Lys Cys Gly Ser
                85                  90                  95

CCG GGC CAG CTC CAG GGC CTC AGC ATC TTT GCA CAG GAC CTG CGC CTC     336
Pro Gly Gln Leu Gln Gly Leu Ser Ile Phe Ala Gln Asp Leu Arg Leu
            100                 105                 110

TGC CTG GAT GAG GCC CTC TCC TGG GAC TGT TTC GCC CTC TCG CTG CTG     384
Cys Leu Asp Glu Ala Leu Ser Trp Asp Cys Phe Ala Leu Ser Leu Leu
        115                 120                 125

GCT GTG GCT CTG GGC CTG GGT GTC CCC ATG CTG CAT CAC CTC TGT GGC     432
Ala Val Ala Leu Gly Leu Gly Val Pro Met Leu His His Leu Cys Gly
    130                 135                 140

TGG GAC CTC TGG TAC TGC TTC CAC CTG TGC CTG GCC TGG CTT CCC TGG     480
Trp Asp Leu Trp Tyr Cys Phe His Leu Cys Leu Ala Trp Leu Pro Trp
145                 150                 155                 160
```

```
CGG GGG CGG CAA AGT GGG CGA GAT GAG GAT GCC CTG CCC TAC GAT GCC      528
Arg Gly Arg Gln Ser Gly Arg Asp Glu Asp Ala Leu Pro Tyr Asp Ala
                165                 170                 175

TTC GTG GTC TTC GAC AAA ACG CAG AGC GCA GTG GCA GAC TGG GTG TAC      576
Phe Val Val Phe Asp Lys Thr Gln Ser Ala Val Ala Asp Trp Val Tyr
            180                 185                 190

AAC GAG CTT CGG GGG CAG CTG GAG GAG TGC CGT GGG CGC TGG GCA CTC      624
Asn Glu Leu Arg Gly Gln Leu Glu Glu Cys Arg Gly Arg Trp Ala Leu
        195                 200                 205

CGC CTG TGC CTG GAG GAA CGC GAC TGG CTG CCT GGC AAA ACC CTC TTT      672
Arg Leu Cys Leu Glu Glu Arg Asp Trp Leu Pro Gly Lys Thr Leu Phe
    210                 215                 220

GAG AAC CTG TGG GCC TCG GTC TAT GGC AGC CGC AAG ACG CTG TTT GTG      720
Glu Asn Leu Trp Ala Ser Val Tyr Gly Ser Arg Lys Thr Leu Phe Val
225                 230                 235                 240

CTG GCC CAC ACG GAC CGG GTC AGT GGT CTC TTG CGC GCC AGC TTC CTG      768
Leu Ala His Thr Asp Arg Val Ser Gly Leu Leu Arg Ala Ser Phe Leu
                245                 250                 255

CTG GCC CAG CAG CGC CTG CTG GAG GAC CGC AAG GAC GTC GTG GTG CTG      816
Leu Ala Gln Gln Arg Leu Leu Glu Asp Arg Lys Asp Val Val Val Leu
            260                 265                 270

GTG ATC CTG AGC CCT GAC GGC CGC CGC TCC CGC TAC GAG CGG CTG CGC      864
Val Ile Leu Ser Pro Asp Gly Arg Arg Ser Arg Tyr Glu Arg Leu Arg
        275                 280                 285

CAG CGC CTC TGC CGC CAG AGT GTC CTC CTC TGG CCC CAC CAG CCC AGT      912
Gln Arg Leu Cys Arg Gln Ser Val Leu Leu Trp Pro His Gln Pro Ser
    290                 295                 300

GGT CAG CGC AGC TTC TGG GCC CAG CTG GGC ATG GCC CTG ACC AGG GAC      960
Gly Gln Arg Ser Phe Trp Ala Gln Leu Gly Met Ala Leu Thr Arg Asp
305                 310                 315                 320

AAC CAC CAC TTC TAT AAC CGG AAC TTC TGC CAG GGA CCC ACG GCC GAA     1008
Asn His His Phe Tyr Asn Arg Asn Phe Cys Gln Gly Pro Thr Ala Glu
                325                 330                 335

TAGCCGTGAG CCGGAATCCT GCACGGTGCC ACCTCCACAC TCACCTCACC TCTGCCTGCC    1068

TGGTCTGACC CTCCCCTGCT CGCCTCCCTC ACCCCACACC TGACACAGAG CAGGCACTCA    1128

ATAAATGCTA CCGAAGGCTA AAAAAAAAAA AAAAAAAAAA AACCA                    1173
```

LPAGTRLRRLDVSCNSISFVAPGFFSKAKELRELNLSANALKTVDHSWFGPLASALQILDVSANPLHCACGAAFM
DFLLEVQAAVPGLPSRVKCGSPGQLQGLSIFAQDLRLCLDEALSWDCFALSLLAVALGLGVPMLHHLCGWDLWYC
FHLCLAWLPWRGRQSGRDEDALPYDAFVVFDKTQSAVADWVYNELRGQLEECRGRWALRLCLEERDWLPGKTLFE
NLWASVYGSRKTLFVLAHTDRVSGLLRASFLLAQQRLLEDRKDVVVLVILSPDGRRSRY.RLRQRLCRQSVLLWP
HQPSGQRSFWAQLGMALTRDNHHFYNRNFCQGPTAE

Further primate, e.g., human, DTLR10 (SEQ ID NO: 42 and 43):

```
atg ccc atg aag tgg agt ggg tgg agg tgg agc tgg ggg ccg gcc act   48
Met Pro Met Lys Trp Ser Gly Trp Arg Trp Ser Trp Gly Pro Ala Thr
            -45                 -40                 -35

cac aca gcc ctc cca ccc cca cag ggt ttc tgc cgc agc gcc ctg cac   96
His Thr Ala Leu Pro Pro Pro Gln Gly Phe Cys Arg Ser Ala Leu His
        -30                 -25                 -20

ccg ctg tct ctc ctg gtg cag gcc atc atg ctg gcc atg acc ctg gcc   144
Pro Leu Ser Leu Leu Val Gln Ala Ile Met Leu Ala Met Thr Leu Ala
    -15                 -10                 -5                  -1

ctg ggt acc ttg cct gcc ttc cta ccc tgt gag ctc cag ccc cac ggc   192
Leu Gly Thr Leu Pro Ala Phe Leu Pro Cys Glu Leu Gln Pro His Gly
  1                 5                   10                  15

ctg gtg aac tgc aac tgg ctg ttc ctg aag tct gtg ccc cac ttc tcc   240
Leu Val Asn Cys Asn Trp Leu Phe Leu Lys Ser Val Pro His Phe Ser
              20                  25                  30

atg gca gca ccc cgt ggc aat gtc acc agc ctt tcc ttg tcc tcc aac   288
Met Ala Ala Pro Arg Gly Asn Val Thr Ser Leu Ser Leu Ser Ser Asn
          35                  40                  45

cgc atc cac cac ctc cat gat tct gac ttt gcc cac ctg ccc agc ctg   336
Arg Ile His His Leu His Asp Ser Asp Phe Ala His Leu Pro Ser Leu
      50                  55                  60

cgg cat ctc aac ctc aag tgg aac tgc ccg ccg gtt ggc ctc agc ccc   384
Arg His Leu Asn Leu Lys Trp Asn Cys Pro Pro Val Gly Leu Ser Pro
  65                  70                  75                  80

atg cac ttc ccc tgc cac atg acc atc gag ccc agc acc ttc ttg gct   432
Met His Phe Pro Cys His Met Thr Ile Glu Pro Ser Thr Phe Leu Ala
                  85                  90                  95

gtg ccc acc ctg gaa gag cta aac ctg agc tac aac aac atc atg act   480
Val Pro Thr Leu Glu Glu Leu Asn Leu Ser Tyr Asn Asn Ile Met Thr
              100                 105                 110

gtg cct gcg ctg ccc aaa tcc ctc ata tcc ctg tcc ctc agc cat acc   528
Val Pro Ala Leu Pro Lys Ser Leu Ile Ser Leu Ser Leu Ser His Thr
          115                 120                 125

aac atc ctg atg cta gac tct gcc agc ctc gcc ggc ctg cat gcc ctg   576
Asn Ile Leu Met Leu Asp Ser Ala Ser Leu Ala Gly Leu His Ala Leu
      130                 135                 140
```

```
cgc ttc cta ttc atg gac ggc aac tgt tat tac aag aac ccc tgc agg   624
Arg Phe Leu Phe Met Asp Gly Asn Cys Tyr Tyr Lys Asn Pro Cys Arg
145                 150                 155                 160

cag gca ctg gag gtg gcc ccg ggt gcc ctc ctt ggc ctg ggc aac ctc   672
Gln Ala Leu Glu Val Ala Pro Gly Ala Leu Leu Gly Leu Gly Asn Leu
                165                 170                 175

acc cac ctg tca ctc aag tac aac aac ctc act gtg gtg ccc cgc aac   720
Thr His Leu Ser Leu Lys Tyr Asn Asn Leu Thr Val Val Pro Arg Asn
            180                 185                 190

ctg cct tcc agc ctg gag tat ctg ctg ttg tcc tac aac cgc atc gtc   768
Leu Pro Ser Ser Leu Glu Tyr Leu Leu Leu Ser Tyr Asn Arg Ile Val
        195                 200                 205

aaa ctg gcg cct gag gac ctg gcc aat ctg acc gcc ctg cgt gtg ctc   816
Lys Leu Ala Pro Glu Asp Leu Ala Asn Leu Thr Ala Leu Arg Val Leu
    210                 215                 220

gat gtg ggc gga aat tgc cgc cgc tgc gac cac gct ccc aac ccc tgc   864
Asp Val Gly Gly Asn Cys Arg Arg Cys Asp His Ala Pro Asn Pro Cys
225                 230                 235                 240

atg gag tgc cct cgt cac ttc ccc cag cta cat ccc gat acc ttc agc   912
Met Glu Cys Pro Arg His Phe Pro Gln Leu His Pro Asp Thr Phe Ser
                245                 250                 255

cac ctg agc cgt ctt gaa ggc ctg gtg ttg aag gac agt tct ctc tcc   960
His Leu Ser Arg Leu Glu Gly Leu Val Leu Lys Asp Ser Ser Leu Ser
            260                 265                 270

tgg ctg aat gcc agt tgg ttc cgt ggg ctg gga aac ctc cga gtg ctg  1008
Trp Leu Asn Ala Ser Trp Phe Arg Gly Leu Gly Asn Leu Arg Val Leu
        275                 280                 285

gac ctg agt gag aac ttc ctc tac aaa tgc atc act aaa acc aag gcc  1056
Asp Leu Ser Glu Asn Phe Leu Tyr Lys Cys Ile Thr Lys Thr Lys Ala
    290                 295                 300

ttc cag ggc cta aca cag ctg cgc aag ctt aac ctg tcc ttc aat tac  1104
Phe Gln Gly Leu Thr Gln Leu Arg Lys Leu Asn Leu Ser Phe Asn Tyr
305                 310                 315                 320

caa aag agg gtg tcc ttt gcc cac ctg tct ctg gcc cct tcc ttc ggg  1152
Gln Lys Arg Val Ser Phe Ala His Leu Ser Leu Ala Pro Ser Phe Gly
                325                 330                 335

agc ctg gtc gcc ctg aag gag ctg gac atg cac ggc atc ttc ttc cgc  1200
Ser Leu Val Ala Leu Lys Glu Leu Asp Met His Gly Ile Phe Phe Arg
            340                 345                 350

tca ctc gat gag acc acg ctc cgg cca ctg gcc cgc ctg ccc atg ctc  1248
Ser Leu Asp Glu Thr Thr Leu Arg Pro Leu Ala Arg Leu Pro Met Leu
        355                 360                 365
```

```
cag act ctg cgt ctg cag atg aac ttc atc aac cag gcc cag ctc ggc   1296
Gln Thr Leu Arg Leu Gln Met Asn Phe Ile Asn Gln Ala Gln Leu Gly
    370                 375                 380

atc ttc agg gcc ttc cct ggc ctg cgc tac gtg gac ctg tcg gac aac   1344
Ile Phe Arg Ala Phe Pro Gly Leu Arg Tyr Val Asp Leu Ser Asp Asn
385                 390                 395                 400

cgc atc agc gga gct tcg gag ctg aca gcc acc atg ggg gag gca gat   1392
Arg Ile Ser Gly Ala Ser Glu Leu Thr Ala Thr Met Gly Glu Ala Asp
                405                 410                 415

gga ggg gag aag gtc tgg ctg cag cct ggg gac ctt gct ccg gcc cca   1440
Gly Gly Glu Lys Val Trp Leu Gln Pro Gly Asp Leu Ala Pro Ala Pro
            420                 425                 430

gtg gac act ccc agc tct gaa gac ttc agg ccc aac tgc agc acc ctc   1488
Val Asp Thr Pro Ser Ser Glu Asp Phe Arg Pro Asn Cys Ser Thr Leu
        435                 440                 445

aac ttc acc ttg gat ctg tca cgg aac aac ctg gtg acc gtg cag ccg   1536
Asn Phe Thr Leu Asp Leu Ser Arg Asn Asn Leu Val Thr Val Gln Pro
    450                 455                 460

gag atg ttt gcc cag ctc tcg cac ctg cag tgc ctg cgc ctg agc cac   1584
Glu Met Phe Ala Gln Leu Ser His Leu Gln Cys Leu Arg Leu Ser His
465                 470                 475                 480

aac tgc atc tcg cag gca gtc aat ggc tcc cag ttc ctg ccg ctg acc   1632
Asn Cys Ile Ser Gln Ala Val Asn Gly Ser Gln Phe Leu Pro Leu Thr
                485                 490                 495

ggt ctg cag gtg cta gac ctg tcc cac aat aag ctg gac ctc tac cac   1680
Gly Leu Gln Val Leu Asp Leu Ser His Asn Lys Leu Asp Leu Tyr His
            500                 505                 510

gag cac tca ttc acg gag cta cca cga ctg gag gcc ctg gac ctc agc   1728
Glu His Ser Phe Thr Glu Leu Pro Arg Leu Glu Ala Leu Asp Leu Ser
        515                 520                 525

tac aac agc cag ccc ttt ggc atg cag ggc gtg ggc cac aac ttc agc   1776
Tyr Asn Ser Gln Pro Phe Gly Met Gln Gly Val Gly His Asn Phe Ser
    530                 535                 540

ttc gtg gct cac ctg cgc acc ctg cgc cac ctc agc ctg gcc cac aac   1824
Phe Val Ala His Leu Arg Thr Leu Arg His Leu Ser Leu Ala His Asn
545                 550                 555                 560

aac atc cac agc caa gtg tcc cag cag ctc tgc agt acg tcg ctg cgg   1872
Asn Ile His Ser Gln Val Ser Gln Gln Leu Cys Ser Thr Ser Leu Arg
                565                 570                 575

gcc ctg gac ttc agc ggc aat gca ctg ggc cat atg tgg gcc gag gga   1920
Ala Leu Asp Phe Ser Gly Asn Ala Leu Gly His Met Trp Ala Glu Gly
            580                 585                 590
```

```
gac ctc tat ctg cac ttc ttc caa ggc ctg agc ggt ttg atc tgg ctg   1968
Asp Leu Tyr Leu His Phe Phe Gln Gly Leu Ser Gly Leu Ile Trp Leu
        595                 600                 605

gac ttg tcc cag aac cgc ctg cac acc ctc ctg ccc caa acc ctg cgc   2016
Asp Leu Ser Gln Asn Arg Leu His Thr Leu Leu Pro Gln Thr Leu Arg
    610                 615                 620

aac ctc ccc aag agc cta cag gtg ctg cgt ctc cgt gac aat tac ctg   2064
Asn Leu Pro Lys Ser Leu Gln Val Leu Arg Leu Arg Asp Asn Tyr Leu
625                 630                 635                 640

gcc ttc ttt aag tgg tgg agc ctc cac ttc ctg ccc aaa ctg gaa gtc   2112
Ala Phe Phe Lys Trp Trp Ser Leu His Phe Leu Pro Lys Leu Glu Val
                645                 650                 655

ctc gac ctg gca gga aac cag ctg aag gcc ctg acc aat ggc agc ctg   2160
Leu Asp Leu Ala Gly Asn Gln Leu Lys Ala Leu Thr Asn Gly Ser Leu
            660                 665                 670

cct gct ggc acc cgg ctc cgg agg ctg gat gtc agc tgc aac agc atc   2208
Pro Ala Gly Thr Arg Leu Arg Arg Leu Asp Val Ser Cys Asn Ser Ile
        675                 680                 685

agc ttc gtg gcc ccc ggc ttc ttt tcc aag gcc aag gag ctg cga gag   2256
Ser Phe Val Ala Pro Gly Phe Phe Ser Lys Ala Lys Glu Leu Arg Glu
    690                 695                 700

ctc aac ctt agc gcc aac gcc ctc aag aca gtg gac cac tcc tgg ttt   2304
Leu Asn Leu Ser Ala Asn Ala Leu Lys Thr Val Asp His Ser Trp Phe
705                 710                 715                 720

ggg ccc ctg gcg agt gcc ctg caa ata cta gat gta agc gcc aac cct   2352
Gly Pro Leu Ala Ser Ala Leu Gln Ile Leu Asp Val Ser Ala Asn Pro
                725                 730                 735

ctg cac tgc gcc tgt ggg gcg gcc ttt atg gac ttc ctg ctg gag gtg   2400
Leu His Cys Ala Cys Gly Ala Ala Phe Met Asp Phe Leu Leu Glu Val
            740                 745                 750

cag gct gcc gtg ccc ggt ctg ccc agc cgg gtg aag tgt ggc agt ccg   2448
Gln Ala Ala Val Pro Gly Leu Pro Ser Arg Val Lys Cys Gly Ser Pro
        755                 760                 765

ggc cag ctc cag ggc ctc agc atc ttt gca cag gac ctg cgc ctc tgc   2496
Gly Gln Leu Gln Gly Leu Ser Ile Phe Ala Gln Asp Leu Arg Leu Cys
    770                 775                 780

ctg gat gag gcc ctc tcc tgg gac tgt ttc gcc ctc tcg ctg ctg gct   2544
Leu Asp Glu Ala Leu Ser Trp Asp Cys Phe Ala Leu Ser Leu Leu Ala
785                 790                 795                 800

gtg gct ctg ggc ctg ggt gtg ccc atg ctg cat cac ctc tgt ggc tgg   2592
Val Ala Leu Gly Leu Gly Val Pro Met Leu His His Leu Cys Gly Trp
                805                 810                 815
```

```
gac ctc tgg tac tgc ttc cac ctg tgc ctg gcc tgg ctt ccc tgg cgg   2640
Asp Leu Trp Tyr Cys Phe His Leu Cys Leu Ala Trp Leu Pro Trp Arg
            820                 825                 830

ggg cgg caa agt ggg cga gat gag gat gcc ctg ccc tac gat gcc ttc   2688
Gly Arg Gln Ser Gly Arg Asp Glu Asp Ala Leu Pro Tyr Asp Ala Phe
        835                 840                 845

gtg gtc ttc gac aaa acg cag agc gca gtg gca gac tgg gtg tac aac   2736
Val Val Phe Asp Lys Thr Gln Ser Ala Val Ala Asp Trp Val Tyr Asn
    850                 855                 860

gag ctt cgg ggg cag ctg gag gag tgc cgt ggg cgc tgg gca ctc cgc   2784
Glu Leu Arg Gly Gln Leu Glu Glu Cys Arg Gly Arg Trp Ala Leu Arg
865                 870                 875                 880

ctg tgc ctg gag gaa cgc gac tgg ctg cct ggc aaa acc ctc ttt gag   2832
Leu Cys Leu Glu Glu Arg Asp Trp Leu Pro Gly Lys Thr Leu Phe Glu
                885                 890                 895

aac ctg tgg gcc tcg gtc tat ggc agc cgc aag acg ctg ttt gtg ctg   2880
Asn Leu Trp Ala Ser Val Tyr Gly Ser Arg Lys Thr Leu Phe Val Leu
            900                 905                 910

gcc cac acg gac cgg gtc agt ggt ctc ttg cgc gcc agc ttc ctg ctg   2928
Ala His Thr Asp Arg Val Ser Gly Leu Leu Arg Ala Ser Phe Leu Leu
        915                 920                 925

gcc cag cag cgc ctg ctg gag gac cgc aag gac gtc gtg gtg ctg gtg   2976
Ala Gln Gln Arg Leu Leu Glu Asp Arg Lys Asp Val Val Val Leu Val
    930                 935                 940

atc ctg agc cct gac ggc cgc cgc tcc cgc tat gtg cgg ctg cgc cag   3024
Ile Leu Ser Pro Asp Gly Arg Arg Ser Arg Tyr Val Arg Leu Arg Gln
945                 950                 955                 960

cgc ctc tgc cgc cag agt gtc ctc ctc tgg ccc cac cag ccc agt ggt   3072
Arg Leu Cys Arg Gln Ser Val Leu Leu Trp Pro His Gln Pro Ser Gly
                965                 970                 975

cag cgc agc ttc tgg gcc cag ctg ggc atg gcc ctg acc agg gac aac   3120
Gln Arg Ser Phe Trp Ala Gln Leu Gly Met Ala Leu Thr Arg Asp Asn
            980                 985                 990

cac cac ttc tat aac cgg aac ttc tgc cag gga ccc acg gcc gaa tag   3168
His His Phe Tyr Asn Arg Asn Phe Cys Gln Gly Pro Thr Ala Glu
        995                1000                1005
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MPMKWSGWRWSWGPATHTALPPPQGFCRSALHPLSLLVQAIMLAMTLALGTLPAFLPCELQPHGLVNCN
WLFLKSVPHFSMAAPRGNVTSLSLSSNRIHHLHDSDFAHLPSLRHLNLKWNCPPVGLSPMHFPCHMTIE
PSTFLAVPTLEELNLSYNNIMTVPALPKSLISLSLSHTNILMLDSASLAGLHALRFLFMDGNCYYKNPC
RQALEVAPGALLGLGNLTHLSLKYNNLTVVPRNLPSSLEYLLLSYNRIVKLAPEDLANLTALRVLDVGG
NCRRCDHAPNPCMECPRHFPQLHPDTFSHLSRLEGLVLKDSSLSWLNASWFRGLGNLRVLDLSENFLYK
CITKTKAFQGLTQLRKLNLSFNYQKRVSFAHLSLAPSFGSLVALKELDMHGIFFRSLDETTLRPLARLP
MLQTLRLQMNFINQAQLGIFRAFPGLRYVDLSDNRISGASELTATMGEADGGEKVWLQPGDLAPAPVDT
PSSEDFRPNCSTLNFTLDLSRNNLVTVQPEMFAQLSHLQCLRLSHNCISQAVNGSQFLPLTGLQVLDLS
HNKLDLYHEHSFTELPRLEALDLSYNSQPFGMQGVGHNFSFVAHLRTLRHLSLAHNNIHSQVSQQLCST
SLRALDFSGNALGHMWAEGDLYLHFFQGLSGLIWLDLSQNRLHTLLPQTLRNLPKSLQVLRLRDNYLAF
FKWWSLHFLPKLEVLDLAGNQLKALTNGSLPAGTRLRRLDVSCNSISFVAPGFFSKAKELRELNLSANA
LKTVDHSWFGPLASALQILDVSANPLHCACGAAFMDFLLEVQAAVPGLPSRVKCGSPGQLQGLSIFAQD
LRLCLDEALSWDCFALSLLAVALGLGVPMLHHLCGWDLWYCFHLCLAWLPWRGRQSGRDEDALPYDAFV
VFDKTQSAVADWVYNELRGQLEECRGRWALRLCLEERDWLPGKTLFENLWASVYGSRKTLFVLAHTDRV
SGLLRASFLLAQQRLLEDRKDVVVLVILSPDGRRSRYVRLRQRLCRQSVLLWPHQPSGQRSFWAQLGMA
LTRDNHHFYNRNFCQGPTAE

partial rodent, e.g., mouse DTLR10 nucleotide sequence (SEQ ID NO: 35):

TGGCCCACAC GGACCGCGTC AGTGGCCTCC TGCGCACCAG CTTCCTGCTG GCTCAGCAGC 60

GCCTGTTGGA AGACCGCAAG GACGTGGTGG TGTTGGTGAT CCTGCGTCCG GATGCCCCAC 120

CGTCCCGCTA TGTGCGACTG CGCCAGCGTC TCTGCCGCCA GAGTGTGCTC TTCTGGCCCC 180

AGCGACCCAA CGGGCAGGGG GGCTTCTGGG CCCAGCTGAG TACAGCCCTG ACTAGGGACA 240

ACCGCCACTT CTATAACCAG AACTTCTGCC GGGGACCTAC AGCAGAATAG CTCAGAGCAA 300

CAGCTGGAAA CAGCTGCATC TTCATGTCTG GTTCCCGAGT TGCTCTGCCT GCCTTGCTCT 360

GTCTTACTAC ACCGCTATTT GGCAAGTGCG CAATATATGC TACCAAGCCA CCAGGCCCAC 420

GGAGCAAAGG TTGGCTGTAA AGGGTAGTTT TCTTCCCATG CATCTTTCAG GAGAGTGAAG 480

ATAGACACCA AACCCAC 497

Further rodent, e.g., mouse, DTLR10 (SEQ ID NO: 44 and 45):

```
aac ctg tcc ttc aat tac cgc aag aag gta tcc ttt gcc cgc ctc cac   48
Asn Leu Ser Phe Asn Tyr Arg Lys Lys Val Ser Phe Ala Arg Leu His
 1               5                  10                  15

ctg gca agt tcc ttt aag aac ctg gtg tca ctg cag gag ctg aac atg   96
Leu Ala Ser Ser Phe Lys Asn Leu Val Ser Leu Gln Glu Leu Asn Met
            20                  25                  30

aac ggc atc ttc ttc cgc ttg ctc aac aag tac acg ctc aga tgg ctg  144
Asn Gly Ile Phe Phe Arg Leu Leu Asn Lys Tyr Thr Leu Arg Trp Leu
        35                  40                  45

gcc gat ctg ccc aaa ctc cac act ctg cat ctt caa atg aac ttc atc  192
Ala Asp Leu Pro Lys Leu His Thr Leu His Leu Gln Met Asn Phe Ile
    50                  55                  60
```

```
aac cag gca cag ctc agc atc ttt ggt acc ttc cga gcc ctt cgc ttt   240
Asn Gln Ala Gln Leu Ser Ile Phe Gly Thr Phe Arg Ala Leu Arg Phe
 65                  70                  75                  80

gtg gac ttg tca gac aat cgc atc agt ggg cct tca acg ctg tca gaa   288
Val Asp Leu Ser Asp Asn Arg Ile Ser Gly Pro Ser Thr Leu Ser Glu
                85                  90                  95

gcc acc cct gaa gag gca gat gat gca gag cag gag gag ctg ttg tct   336
Ala Thr Pro Glu Glu Ala Asp Asp Ala Glu Gln Glu Glu Leu Leu Ser
            100                 105                 110

gcg gat cct cac cca gct ccg ctg agc acc cct gct tct aag aac ttc   384
Ala Asp Pro His Pro Ala Pro Leu Ser Thr Pro Ala Ser Lys Asn Phe
        115                 120                 125

atg gac agg tgt aag aac ttc aag ttc aac atg gac ctg tct cgg aac   432
Met Asp Arg Cys Lys Asn Phe Lys Phe Asn Met Asp Leu Ser Arg Asn
    130                 135                 140

aac ctg gtg act atc aca gca gag atg ttt gta aat ctc tca cgc ctc   480
Asn Leu Val Thr Ile Thr Ala Glu Met Phe Val Asn Leu Ser Arg Leu
145                 150                 155                 160

cag tgt ctt agc ctg agc cac aac tca att gca cag gct gtc aat ggc   528
Gln Cys Leu Ser Leu Ser His Asn Ser Ile Ala Gln Ala Val Asn Gly
                165                 170                 175

tct cag ttc ctg ccg ctg acc ggt ctg cag gtg cta gac ctg tcc cac   576
Ser Gln Phe Leu Pro Leu Thr Gly Leu Gln Val Leu Asp Leu Ser His
            180                 185                 190

aat aag ctg gac ctc tac cac gag cac tca ttc acg gag cta cca cga   624
Asn Lys Leu Asp Leu Tyr His Glu His Ser Phe Thr Glu Leu Pro Arg
        195                 200                 205

ctg gag gcc ctg gac ctc agc tac aac agc cag ccc ttt agc atg aag   672
Leu Glu Ala Leu Asp Leu Ser Tyr Asn Ser Gln Pro Phe Ser Met Lys
    210                 215                 220

ggt ata ggc cac aat ttc agt ttt gtg acc cat ctg tcc atg cta cag   720
Gly Ile Gly His Asn Phe Ser Phe Val Thr His Leu Ser Met Leu Gln
225                 230                 235                 240

agc ctt agc ctg gca cac aat gac att cat acc cgt gtg tcc tca cat   768
Ser Leu Ser Leu Ala His Asn Asp Ile His Thr Arg Val Ser Ser His
                245                 250                 255

ctc aac agc aac tca gtg agg ttt ctt gac ttc agc ggc aac ggt atg   816
Leu Asn Ser Asn Ser Val Arg Phe Leu Asp Phe Ser Gly Asn Gly Met
            260                 265                 270

ggc cgc atg tgg gat gag ggg ggc ctt tat ctc cat ttc ttc caa ggc   864
Gly Arg Met Trp Asp Glu Gly Gly Leu Tyr Leu His Phe Phe Gln Gly
        275                 280                 285
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ctg agt ggc gtg ctg aag ctg gac ctg tct caa aat aac ctg cat atc   912
Leu Ser Gly Val Leu Lys Leu Asp Leu Ser Gln Asn Asn Leu His Ile
    290                 295                 300

ctc cgg ccc cag aac ctt gac aac ctc ccc aag agc ctg aag ctg ctg   960
Leu Arg Pro Gln Asn Leu Asp Asn Leu Pro Lys Ser Leu Lys Leu Leu
305                 310                 315                 320

agc ctc cga gac aac tac cta tct ttc ttt aac tgg acc agt ctg tcc  1008
Ser Leu Arg Asp Asn Tyr Leu Ser Phe Phe Asn Trp Thr Ser Leu Ser
                325                 330                 335

ttc cta ccc aac ctg gaa gtc cta gac ctg gca ggc aac cag cta aag  1056
Phe Leu Pro Asn Leu Glu Val Leu Asp Leu Ala Gly Asn Gln Leu Lys
            340                 345                 350

gcc ctg acc aat ggc acc ctg cct aat ggc acc ctc ctc cag aaa ctc  1104
Ala Leu Thr Asn Gly Thr Leu Pro Asn Gly Thr Leu Leu Gln Lys Leu
        355                 360                 365

gat gtc agt agc aac agt atc gtc tct gtg gcc ccc ggc ttc ttt tcc  1152
Asp Val Ser Ser Asn Ser Ile Val Ser Val Ala Pro Gly Phe Phe Ser
    370                 375                 380

aag gcc aag gag ctg cga gag ctc aac ctt agc gcc aac gcc ctc aag  1200
Lys Ala Lys Glu Leu Arg Glu Leu Asn Leu Ser Ala Asn Ala Leu Lys
385                 390                 395                 400

aca gtg gac cac tcc tgg ttt ggg ccc att gtg atg aac ctg aca gtt  1248
Thr Val Asp His Ser Trp Phe Gly Pro Ile Val Met Asn Leu Thr Val
                405                 410                 415

cta gac gtg aga agc aac cct ctg cac tgt gcc tgt ggg gca gcc ttc  1296
Leu Asp Val Arg Ser Asn Pro Leu His Cys Ala Cys Gly Ala Ala Phe
            420                 425                 430

gta gac tta ctg ttg gag gtg cag acc aag gtg cct ggc ctg gct aat  1344
Val Asp Leu Leu Leu Glu Val Gln Thr Lys Val Pro Gly Leu Ala Asn
        435                 440                 445

ggt gtg aag tgt ggc agc ccc ggc cag ctg cag ggc cgt agc atc ttc  1392
Gly Val Lys Cys Gly Ser Pro Gly Gln Leu Gln Gly Arg Ser Ile Phe
    450                 455                 460

gcg cag gac ctg cgg ctg tgc ctg gat gag gtc ctc tct tgg gac tgc  1440
Ala Gln Asp Leu Arg Leu Cys Leu Asp Glu Val Leu Ser Trp Asp Cys
465                 470                 475                 480

ttt ggc ctt tca ctc ttg gct gtg gcc gtg ggc atg gtg gtg cct ata  1488
Phe Gly Leu Ser Leu Leu Ala Val Ala Val Gly Met Val Val Pro Ile
                485                 490                 495

ctg cac cat ctc tgc ggc tgg gac gtc tgg tac tgt ttt cat ctg tgc  1536
Leu His His Leu Cys Gly Trp Asp Val Trp Tyr Cys Phe His Leu Cys
            500                 505                 510
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ctg gca tgg cta cct ttg cta gcc cgc agc cga cgc agc gcc caa act   1584
Leu Ala Trp Leu Pro Leu Leu Ala Arg Ser Arg Arg Ser Ala Gln Thr
        515                 520                 525

ctc cct tat gat gcc ttc gtg gtg ttc gat aag gca cag agc gca gtt   1632
Leu Pro Tyr Asp Ala Phe Val Val Phe Asp Lys Ala Gln Ser Ala Val
    530                 535                 540

gcc gac tgg gtg tat aac gag ctg cgg gtg cgg ctg gag gag cgg cgc   1680
Ala Asp Trp Val Tyr Asn Glu Leu Arg Val Arg Leu Glu Glu Arg Arg
545                 550                 555                 560

ggc cgc tgg gca ctc cgc ctg tgc ctg gag gac cga gat tgg ctg cct   1728
Gly Arg Trp Ala Leu Arg Leu Cys Leu Glu Asp Arg Asp Trp Leu Pro
                565                 570                 575

ggc cag acg ctc ttc gag aac ctc tgg gct tcc atc tat ggg agc cgc   1776
Gly Gln Thr Leu Phe Glu Asn Leu Trp Ala Ser Ile Tyr Gly Ser Arg
            580                 585                 590

aag act cta ttt gtg ctg gcc cac acg gac cgc gtc agt cgc ctc ctg   1824
Lys Thr Leu Phe Val Leu Ala His Thr Asp Arg Val Ser Gly Leu Leu
        595                 600                 605

cgc acc agc ttc ctg ctg gct cag cag cgc ctg ttg gaa gac cgc aag   1872
Arg Thr Ser Phe Leu Leu Ala Gln Gln Arg Leu Leu Glu Asp Arg Lys
    610                 615                 620

gac gtg gtg gtg ttg gtg atc ctg cgt ccg gat gcc cac cgc tcc cgc   1920
Asp Val Val Val Leu Val Ile Leu Arg Pro Asp Ala His Arg Ser Arg
625                 630                 635                 640

tat gtg cga ctg cgc cag cgt ctc tgc cgc cag agt gtg ctc ttc tgg   1968
Tyr Val Arg Leu Arg Gln Arg Leu Cys Arg Gln Ser Val Leu Phe Trp
                645                 650                 655

ccc cag cag ccc aac ggg cag ggg ggc ttc tgg gcc cag ctg agt aca   2016
Pro Gln Gln Pro Asn Gly Gln Gly Gly Phe Trp Ala Gln Leu Ser Thr
            660                 665                 670

gcc ctg act agg gac aac cgc cac ttc tat aac cag aac ttc tgc cgg   2064
Ala Leu Thr Arg Asp Asn Arg His Phe Tyr Asn Gln Asn Phe Cys Arg
        675                 680                 685

gga cct aca gca gaa tagctcagag caacagctgg aaacagctgc atcttcatgt   2119
Gly Pro Thr Ala Glu
    690

ctggttcccg agttgctctg cctgccttgc tctgtcttac tacaccgcta tttggcaagt 2179

gcgcaatata tgctaccaag ccaccaggcc cacggagcaa aggttggctg taaagggtag 2239

ttttcttccc atgcatcttt caggagagtg aagatagaca ccaaacccac           2289
```

NLSFNYRKKVSFARLHLASSFKNLVSLQELNMNGIFFRLLNKYTLRWLADLPKLHTLHLQMNFINQAQL
SIFGTFRALRFVDLSDNRISGPSTLSEATPEEADDAEQEELLSADPHPAPLSTPASKNFMDRCKNFKFN
MDLSRNNLVTITAEMFVNLSRLQCLSLSHNSIAQAVNGSQFLPLTGLQVLDLSHNKLDLYHEHSFTELP
RLEALDLSYNSQPFSMKGIGHNFSFVTHLSMLQSLSLAHNDIHTRVSSHLNSNSVRFLDFSGNGMGRMW
DEGGLYLHFFQGLSGVLKLDLSQNNLHILRPQNLDNLPKSLKLLSLRDNYLSFFNWTSLSFLPNLEVLD
LAGNQLKALTNGTLPNGTLLQKLDVSSNSIVSVAPGFFSKAKELRELNLSANALKTVDHSWFGPIVMNL
TVLDVRSNPLHCACGAAFVDLLLEVQTKVPGLANGVKCGSPGQLQGRSIFAQDLRLCLDEVLSWDCFGL
SLLAVAVGMVVPILHHLCGWDVWYCFHLCLAWLPLLARSRRSAQTLPYDAFVVFDKAQSAVADWVYNEL
RVRLEERRGRWALRLCLEDRDWLPGQTLFENLWASIYGSRKTLFVLAHTDRVSGLLRTSFLLAQQRLLE
DRKDVVVLVILRPDAHRSRYVRLRQRLCRQSVLFWPQQPNGQGGFWAQLSTALTRDNRHFYNQNFCRGP
TAE

Table 11: Comparison of intracellular domains of human DTLRs. DTLR1 is SEQ ID NO: 2; DTLR2 is SEQ ID NO: 4; DTLR3 is SEQ ID NO: 6; DTLR4 is SEQ ID NO: 8; DTLR5 is SEQ ID NO: 10; and DTLR6 is SEQ ID NO: 12. Particularly important and conserved, e.g., characteristic, residues correspond, across the DTLRs, to SEQ ID NO: 18 residues tyr10-tyr13; trp26; cys46; trp52; pro54-gly55; ser69; lys71; trp134-pro135; and phe144-trp145.

```
DTLR1    QRNLQFHAFISYSGHD---SFWVKNELLPNLEKEG-----MQICLHERNF
DTLR9    KENLQFHAFISYSEHD---SAWVKSELVPYLEKED-----IQICLHERNF
DTLR8    ------------------------NELIPNLEKEDGS---ILICLYESYF
DTLR2    SRNICYDAFVSYSERD---AYWVENLMVQELENFNPP---FKLCLHKRDF
DTLR6    SPDCCYDAFIVYDTKDPAVTEWVLAELVAKLEDPREK--HFNLCLEERDW
DTLR7    TSQTFYDAYISYDTKDASVTDWVINELRYHLEESRDK--NVLLCLEERDW
DTLR10   EDALPYDAFVVFDKTXSAVADWVYNELRGQLEECRGRW-ALRLCLEERDW
DTLR4    RGENIYDAFVIYSSQD---EDWVRNELVKNLEEGVPP---FQLCLHYRDF
DTLR5    PDMYKYDAYLCFSSKD---FTWVQNALLKHLDTQYSDQNRFNLCFEERDF
DTLR3    TEQFEYAAYIIHAYKD---KDWVWEHFSSMEKEDQS----LKFCLEERDF
                                   :    .        . :*:    :

DTLR1    VPGKSIVENIITC-IEKSYKSIFVLSPNFVQSEWCH-YELYFAHHNLFHE
DTLR9    VPGKSIVENIINC-IEKSYKSIFVLSPNFVQSEWCH-YELYFAHHNLFHE
DTLR8    DPGKSISENIVSF-IEKSYKSIFVLSPNFVQNEWCH-YEFYFAHHNLFHE
DTLR2    IPGKWIIDNIIDS-IEKSHKTVFVLSENFVKSEWCK-YELDFSHFRLFEE
DTLR6    LPGQPVLENLSQS-IQLSKKTVFVMTDKYAKTENFK-IAFYLSHQRLMDE
DTLR7    DPGLAIIDNLMQS-INQSKKTVFVLTKKYAKSWNFK-TAFYLXLQRLMGE
DTLR10   LPGKTLFENLWAS-VYGSRKTLFVLAHTDRVSGLLR-AIFLLAQQRLLE-
DTLR4    IPGVAIAANIIHEGFHKSRKVIVVVSQHFIQSRWCI-FEYEIAQTWQFLS
DTLR5    VPGENRIANIQDA-IWNSRKIVCLVSRHFLRDGWCL-EAFSYAQGRCLSD
DTLR3    EAGVFELEAIVNS-IKRSRKIIFVITHHLLKDPLCKRFKVHHAVQQAIEQ
          .*      :    .  * * : :::                    :

DTLR1    GSNSLILILLEPIPQYSIPSSYHKLKSLMARRTYLEWPKEKSKRGLFWAN
DTLR9    GSNNLILILLEPIPQNSIPNKYHKLKALMTQRTYLQWPKEKSKRGLFWA-
DTLR8    NSDHIILILLEPIPFYCIPTRYHKLEALLEKKAYLEWPKDRRKCGLFWAN
DTLR2    NNDAAILILLEPIEKKAIPQRFCKLRKIMNTKTYLEWPMDEAQREGFWVN
DTLR6    KVDVIILILFLEKPFQK---SKFLQLRKRLCGSSVLEWPTNPQAHPYFWQC
DTLR7    NMDVIIFILLEPVLQH---SPYLRLRQRICKSSILQWPDNPKAERLFWQT
DTLR10   --------------------------------------------------
DTLR4    SRAGIIFIVLQKVEKT-LLRQQVELYRLLSRNTYLEWEDSVLGRHIFWRR
DTLR5    LNSALIMVVVGSLSQY-QLMKHQSIRGFVQKQQYLRWPEDLQDVGWFLHK
DTLR3    NLDSIILVFLEEIPDYKLNHALCLRRGMFKSHCILNWPVQKERIGAFRHK


DTLR1    LRAAINIKLTEQAKK-------------------------
DTLR9    ----------------------------------------
DTLR8    LRAAVNVNVLATREMYELQTFTELNEESRGSTISLMRTDCL
DTLR2    LRAAIKS---------------------------------
DTLR6    LKNALATDNHVAYSQVFKETV-------------------
DTLR7    LXNVVLTENDSRYNNMYVDSIKQY----------------
DTLR10   ----------------------------------------
DTLR4    LRKALLDGKSWNPEGTVGTGCNWQEATSI-----------
DTLR5    LSQQILKKEKEKKKDNNIPLQTVATIS-------------
DTLR3    LQVALGSKNSVH-----------------------------
```

Transmembrane segments correspond approximately to 802-818 (791-823) of primate DTLR7 SEQ ID NO: 37; 559-575 (550-586) of DTLR8 SEQ ID NO: 39; 553-569 (549-582) of DTLR9 SEQ ID NO: 41; 796-810 (790-814) of DTLR10 SEQ ID NO: 43; and 481-497 (475-503) of DTLR10 SEQ ID NO: 45.

As used herein, the term DNAX Toll like receptor 2 (DTLR2) shall be used to describe a protein comprising a protein or peptide segment having or sharing the amino acid sequence shown in Table 2, or a substantial fragment thereof. Similarly, with a DTLR3 and Table 3; DTLR4 and Table 4; DTLR5 and Table 5; DTLR6 and Table 6; DTLR7 and Table 7; DTLR8 and Table 8; DTLR9 and Table 9; and DTLR10 and Table 10. Rodent, e.g., mouse, DTLR11 sequence is provided, e.g., in EST AA739083; DTLR13 in ESTAI019567; DTLR14 in ESTs AI390330 and AA244663.

The invention also includes a protein variations of the respective DTLR allele whose sequence is provided, e.g., a mutein agonist or antagonist. Typically, such agonists or antagonists will exhibit less than about 10% sequence differences, and thus will often have between 1- and 11-fold substitutions, e.g., 2-, 3-, 5-, 7-fold, and others. It also encompasses allelic and other variants, e.g., natural polymorphic, of the protein described. Typically, it will bind to its corresponding biological receptor with high affinity, e.g., at least about 100 nM, usually better than about 30 nM, preferably better than about 10 nM, and more preferably at better than about 3 nM. The term shall also be used herein to refer to related naturally occurring forms, e.g., alleles, polymorphic variants, and metabolic variants of the mammalian protein.

This invention also encompasses proteins or peptides having substantial amino acid sequence identity with the amino acid sequence in Table 2. It will include sequence variants with relatively few substitutions, e.g., preferably less than about 3-5. Similar features apply to

the other DTLR sequences provided in Tables 3, 4, 5, 6, 7, 8, 9, or 10.

A substantial polypeptide "fragment", or "segment", is a stretch of amino acid residues of at least about 8 amino acids, generally at least 10 amino acids, more generally at least 12 amino acids, often at least 14 amino acids, more often at least 16 amino acids, typically at least 18 amino acids, more typically at least 20 amino acids, usually at least 22 amino acids, more usually at least 24 amino acids, preferably at least 26 amino acids, more preferably at least 28 amino acids, and, in particularly preferred embodiments, at least about 30 or more amino acids. Sequences of segments of different proteins can be compared to one another over appropriate length stretches.

Amino acid sequence homology, or sequence identity, is determined by optimizing residue matches, if necessary, by introducing gaps as required. See, e.g., Needleham, et al., (1970) J. Mol. Biol. 48:443-453; Sankoff, et al., (1983) chapter one in Time Warps, String Edits, and Macromolecules: The Theory and Practice of Sequence Comparison, Addison-Wesley, Reading, MA; and software packages from IntelliGenetics, Mountain View, CA; the University of Wisconsin Genetics Computer Group (GCG), Madison, WI; and the NCBI (NIH); each of which is incorporated herein by reference. This changes when considering conservative substitutions as matches. Conservative substitutions typically include substitutions within the following groups: glycine, alanine; valine, isoleucine, leucine; aspartic acid, glutamic acid; asparagine, glutamine; serine, threonine; lysine, arginine; and phenylalanine, tyrosine. Homologous amino acid sequences are intended to include natural allelic and interspecies variations in the cytokine sequence. Typical homologous proteins or peptides will have from 50-100% homology (if gaps can be introduced), to 60-100% homology

(if conservative substitutions are included) with an amino acid sequence segment of Table 2, 3, 4, 5, 6, 7, 8, 9, or 10. Homology measures will be at least about 70%, generally at least 76%, more generally at least 81%, often at least 85%, more often at least 88%, typically at least 90%, more typically at least 92%, usually at least 94%, more usually at least 95%, preferably at least 96%, and more preferably at least 97%, and in particularly preferred embodiments, at least 98% or more. The degree of homology will vary with the length of the compared segments. Homologous proteins or peptides, such as the allelic variants, will share most biological activities with the embodiments described in Table 2, 3, 4, 6, 7, 8, 9, or 10. Particularly interesting regions of comparison, at the amino acid or nucleotide levels, correspond to those within each of the blocks 1-10, or intrablock regions, corresponding to those indicated in Figures 2A-2B.

As used herein, the term "biological activity" is used to describe, without limitation, effects on inflammatory responses, innate immunity, and/or morphogenic development by respective ligands. For example, these receptors should, like IL-1 receptors, mediate phosphatase or phosphorylase activities, which activities are easily measured by standard procedures. See, e.g., Hardie, et al. (eds. 1995) The Protein Kinase FactBook vols. I and II, Academic Press, San Diego, CA; Hanks, et al. (1991) Meth. Enzymol. 200:38-62; Hunter, et al. (1992) Cell 70:375-388; Lewin (1990) Cell 61:743-752; Pines, et al. (1991) Cold Spring Harbor Symp. Quant. Biol. 56:449-463; and Parker, et al. (1993) Nature 363:736-738. The receptors exhibit biological activities much like regulatable enzymes, regulated by ligand binding. However, the enzyme turnover number is more close to an enzyme than a receptor complex. Moreover, the numbers of occupied receptors necessary to induce such enzymatic

activity is less than most receptor systems, and may number closer to dozens per cell, in contrast to most receptors which will trigger at numbers in the thousands per cell. The receptors, or portions thereof, may be useful as phosphate labeling enzymes to label general or specific substrates.

The terms ligand, agonist, antagonist, and analog of, e.g., a DTLR, include molecules that modulate the characteristic cellular responses to Toll ligand like proteins, as well as molecules possessing the more standard structural binding competition features of ligand-receptor interactions, e.g., where the receptor is a natural receptor or an antibody. The cellular responses likely are mediated through binding of various Toll ligands to cellular receptors related to, but possibly distinct from, the type I or type II IL-1 receptors. See, e.g., Belvin and Anderson (1996) Ann. Rev. Cell Dev. Biol. 12:393-416; Morisato and Anderson (1995) Ann. Rev. Genetics 29:371-3991 and Hultmark (1994) Nature 367:116-117.

Also, a ligand is a molecule which serves either as a natural ligand to which said receptor, or an analog thereof, binds, or a molecule which is a functional analog of the natural ligand. The functional analog may be a ligand with structural modifications, or may be a wholly unrelated molecule which has a molecular shape which interacts with the appropriate ligand binding determinants. The ligands may serve as agonists or antagonists, see, e.g., Goodman, et al. (eds. 1990) Goodman & Gilman's: The Pharmacological Bases of Therapeutics, Pergamon Press, New York.

Rational drug design may also be based upon structural studies of the molecular shapes of a receptor or antibody and other effectors or ligands. Effectors may be other proteins which mediate other functions in response to ligand binding, or other proteins which

normally interact with the receptor. One means for determining which sites interact with specific other proteins is a physical structure determination, e.g., x-ray crystallography or 2 dimensional NMR techniques. These will provide guidance as to which amino acid residues form molecular contact regions. For a detailed description of protein structural determination, see, e.g., Blundell and Johnson (1976) Protein Crystallography, Academic Press, New York, which is hereby incorporated herein by reference.

II. Activities

The Toll like receptor proteins will have a number of different biological activities, e.g., in phosphate metabolism, being added to or removed from specific substrates, typically proteins. Such will generally result in modulation of an inflammatory function, other innate immunity response, or a morphological effect. The DTLR2, 3, 4, 5, 6, 7, 8, 9, or 10 proteins are homologous to other Toll like receptor proteins, but each have structural differences. For example, a human DTLR2 gene coding sequence probably has about 70% identity with the nucleotide coding sequence of mouse DTLR2. At the amino acid level, there is also likely to be reasonable identity.

The biological activities of the DTLRs will be related to addition or removal of phosphate moieties to substrates, typically in a specific manner, but occasionally in a non specific manner. Substrates may be identified, or conditions for enzymatic activity may be assayed by standard methods, e.g., as described in Hardie, et al. (eds. 1995) The Protein Kinase FactBook vols. I and II, Academic Press, San Diego, CA; Hanks, et al. (1991) Meth. Enzymol. 200:38-62; Hunter, et al. (1992) Cell 70:375-388; Lewin (1990) Cell 61:743-752; Pines, et al.

(1991) Cold Spring Harbor Symp. Quant. Biol. 56:449-463; and Parker, et al. (1993) Nature 363:736-738.

III. Nucleic Acids

This invention contemplates use of isolated nucleic acid or fragments, e.g., which encode these or closely related proteins, or fragments thereof, e.g., to encode a corresponding polypeptide, preferably one which is biologically active. In addition, this invention covers isolated or recombinant DNA which encodes such proteins or polypeptides having characteristic sequences of the respective DTLRs, individually or as a group. Typically, the nucleic acid is capable of hybridizing, under appropriate conditions, with a nucleic acid sequence segment shown in Tables 2-10, but preferably not with a corresponding segment of Table 1. Said biologically active protein or polypeptide can be a full length protein, or fragment, and will typically have a segment of amino acid sequence highly homologous to one shown in Tables 2-10. Further, this invention covers the use of isolated or recombinant nucleic acid, or fragments thereof, which encode proteins having fragments which are equivalent to the DTLR2-10 proteins. The isolated nucleic acids can have the respective regulatory sequences in the 5' and 3' flanks, e.g., promoters, enhancers, poly-A addition signals, and others from the natural gene.

An "isolated" nucleic acid is a nucleic acid, e.g., an RNA, DNA, or a mixed polymer, which is substantially pure, e.g., separated from other components which naturally accompany a native sequence, such as ribosomes, polymerases, and flanking genomic sequences from the originating species. The term embraces a nucleic acid sequence which has been removed from its naturally occurring environment, and includes recombinant or cloned DNA isolates, which are thereby distinguishable from naturally occurring compositions, and chemically

synthesized analogs or analogs biologically synthesized by heterologous systems. A substantially pure molecule includes isolated forms of the molecule, either completely or substantially pure.

An isolated nucleic acid will generally be a homogeneous composition of molecules, but will, in some embodiments, contain heterogeneity, preferably minor. This heterogeneity is typically found at the polymer ends or portions not critical to a desired biological function or activity.

A "recombinant" nucleic acid is typically defined either by its method of production or its structure. In reference to its method of production, e.g., a product made by a process, the process is use of recombinant nucleic acid techniques, e.g., involving human intervention in the nucleotide sequence. Typically this intervention involves in vitro manipulation, although under certain circumstances it may involve more classical animal breeding techniques. Alternatively, it can be a nucleic acid made by generating a sequence comprising fusion of two fragments which are not naturally contiguous to each other, but is meant to exclude products of nature, e.g., naturally occurring mutants as found in their natural state. Thus, for example, products made by transforming cells with any unnaturally occurring vector is encompassed, as are nucleic acids comprising sequence derived using any synthetic oligonucleotide process. Such a process is often done to replace a codon with a redundant codon encoding the same or a conservative amino acid, while typically introducing or removing a restriction enzyme sequence recognition site. Alternatively, the process is performed to join together nucleic acid segments of desired functions to generate a single genetic entity comprising a desired combination of functions not found in the commonly available natural forms, e.g., encoding a fusion protein. Restriction

enzyme recognition sites are often the target of such artificial manipulations, but other site specific targets, e.g., promoters, DNA replication sites, regulation sequences, control sequences, or other useful features may be incorporated by design. A similar concept is intended for a recombinant, e.g., fusion, polypeptide. This will include a dimeric repeat. Specifically included are synthetic nucleic acids which, by genetic code redundancy, encode equivalent polypeptides to fragments of DTLR2-5 and fusions of sequences from various different related molecules, e.g., other IL-1 receptor family members.

A "fragment" in a nucleic acid context is a contiguous segment of at least about 17 nucleotides, generally at least 21 nucleotides, more generally at least 25 nucleotides, ordinarily at least 30 nucleotides, more ordinarily at least 35 nucleotides, often at least 39 nucleotides, more often at least 45 nucleotides, typically at least 50 nucleotides, more typically at least 55 nucleotides, usually at least 60 nucleotides, more usually at least 66 nucleotides, preferably at least 72 nucleotides, more preferably at least 79 nucleotides, and in particularly preferred embodiments will be at least 85 or more nucleotides. Typically, fragments of different genetic sequences can be compared to one another over appropriate length stretches, particularly defined segments such as the domains described below.

A nucleic acid which codes for a DTLR2-10 will be particularly useful to identify genes, mRNA, and cDNA species which code for itself or closely related proteins, as well as DNAs which code for polymorphic, allelic, or other genetic variants, e.g., from different individuals or related species. Preferred probes for such screens are those regions of the interleukin which are conserved between different polymorphic variants or which contain nucleotides which lack specificity, and will preferably be full length or nearly so. In other situations,

polymorphic variant specific sequences will be more useful.

This invention further covers recombinant nucleic acid molecules and fragments having a nucleic acid sequence identical to or highly homologous to the isolated DNA set forth herein. In particular, the sequences will often be operably linked to DNA segments which control transcription, translation, and DNA replication. These additional segments typically assist in expression of the desired nucleic acid segment.

Homologous, or highly identical, nucleic acid sequences, when compared to one another or Table 2-10 sequences, exhibit significant similarity. The standards for homology in nucleic acids are either measures for homology generally used in the art by sequence comparison or based upon hybridization conditions. Comparative hybridization conditions are described in greater detail below.

Substantial identity in the nucleic acid sequence comparison context means either that the segments, or their complementary strands, when compared, are identical when optimally aligned, with appropriate nucleotide insertions or deletions, in at least about 60% of the nucleotides, generally at least 66%, ordinarily at least 71%, often at least 76%, more often at least 80%, usually at least 84%, more usually at least 88%, typically at least 91%, more typically at least about 93%, preferably at least about 95%, more preferably at least about 96 to 98% or more, and in particular embodiments, as high at about 99% or more of the nucleotides, including, e.g., segments encoding structural domains such as the segments described below. Alternatively, substantial identity will exist when the segments will hybridize under selective hybridization conditions, to a strand or its complement, typically using a sequence derived from Tables 2-10. Typically, selective hybridization will occur when there

is at least about 55% homology over a stretch of at least about 14 nucleotides, more typically at least about 65%, preferably at least about 75%, and more preferably at least about 90%. See, Kanehisa (1984) Nucl. Acids Res. 12:203-213, which is incorporated herein by reference. The length of homology comparison, as described, may be over longer stretches, and in certain embodiments will be over a stretch of at least about 17 nucleotides, generally at least about 20 nucleotides, ordinarily at least about 24 nucleotides, usually at least about 28 nucleotides, typically at least about 32 nucleotides, more typically at least about 40 nucleotides, preferably at least about 50 nucleotides, and more preferably at least about 75 to 100 or more nucleotides.

Stringent conditions, in referring to homology in the hybridization context, will be stringent combined conditions of salt, temperature, organic solvents, and other parameters typically controlled in hybridization reactions. Stringent temperature conditions will usually include temperatures in excess of about 30° C, more usually in excess of about 37° C, typically in excess of about 45° C, more typically in excess of about 55° C, preferably in excess of about 65° C, and more preferably in excess of about 70° C. Stringent salt conditions will ordinarily be less than about 500 mM, usually less than about 400 mM, more usually less than about 300 mM, typically less than about 200 mM, preferably less than about 100 mM, and more preferably less than about 80 mM, even down to less than about 20 mM. However, the combination of parameters is much more important than the measure of any single parameter. See, e.g., Wetmur and Davidson (1968) J. Mol. Biol. 31:349-370, which is hereby incorporated herein by reference.

Alternatively, for sequence comparison, typically one sequence acts as a reference sequence, to which test sequences are compared. When using a sequence comparison

algorithm, test and reference sequences are input into a computer, subsequence coordinates are designated, if necessary, and sequence algorithm program parameters are designated. The sequence comparison algorithm then calculates the percent sequence identity for the test sequence(s) relative to the reference sequence, based on the designated program parameters.

Optical alignment of sequences for comparison can be conducted, e.g., by the local homology algorithm of Smith and Waterman (1981) Adv. Appl. Math. 2:482, by the homology alignment algorithm of Needlman and Wunsch (1970) J. Mol. Biol. 48:443, by the search for similarity method of Pearson and Lipman (1988) Proc. Nat'l Acad. Sci. USA 85:2444, by computerized implementations of these algorithms (GAP, BESTFIT, FASTA, and TFASTA in the Wisconsin Genetics Software Package, Genetics Computer Group, 575 Science Dr., Madison, WI), or by visual inspection (see generally Ausubel et al., supra).

One example of a useful algorithm is PILEUP. PILEUP creates a multiple sequence alignment from a group of related sequences using progressive, pairwise alignments to show relationship and percent sequence identity. It also plots a tree or dendrogram showing the clustering relationships used to create the alignment. PILEUP uses a simplification of the progressive alignment method of Feng and Doolittle (1987) J. Mol. Evol. 35:351-360. The method used is similar to the method described by Higgins and Sharp (1989) CABIOS 5:151-153. The program can align up to 300 sequences, each of a maximum length of 5,000 nucleotides or amino acids. The multiple alignment procedure begins with the pairwise alignment of the two most similar sequences, producing a cluster of two aligned sequences. This cluster is then aligned to the next most related sequence or cluster of aligned sequences. Two clusters of sequences are aligned by a simple extension of the pairwise alignment of two individual sequences. The

final alignment is achieved by a series of progressive, pairwise alignments. The program is run by designating specific sequences and their amino acid or nucleotide coordinates for regions of sequence comparison and by designating the program parameters. For example, a reference sequence can be compared to other test sequences to determine the percent sequence identity relationship using the following parameters: default gap weight (3.00), default gap length weight (0.10), and weighted end gaps.

Another example of algorithm that is suitable for determining percent sequence identity and sequence similarity is the BLAST algorithm, which is described Altschul, et al. (1990) J. Mol. Biol. 215:403-410. Software for performing BLAST analyses is publicly available through the National Center for Biotechnology Information (http:www.ncbi.nlm.nih.gov/). This algorithm involves first identifying high scoring sequence pairs (HSPs) by identifying short words of length W in the query sequence, which either match or satisfy some positive-valued threshold score T when aligned with a word of the same length in a database sequence. T is referred to as the neighborhood word score threshold (Altschul, et al., supra). These initial neighborhood word hits act as seeds for initiating searches to find longer HSPs containing them. The word hits are then extended in both directions along each sequence for as far as the cumulative alignment score can be increased. Extension of the word hits in each direction are halted when: the cumulative alignment score falls off by the quantity X from its maximum achieved value; the cumulative score goes to zero or below, due to the accumulation of one or more negative-scoring residue alignments; or the end of either sequence is reached. The BLAST algorithm parameters W, T, and X determine the sensitivity and speed of the alignment. The BLAST program uses as defaults a wordlength (W) of 11, the BLOSUM62 scoring matrix (see Henikoff and Henikoff (1989)

Proc. Nat'l Acad. Sci. USA 89:10915) alignments (B) of 50, expectation (E) of 10, M=5, N=4, and a comparison of both strands.

In addition to calculating percent sequence identity, the BLAST algorithm also performs a statistical analysis of the similarity between two sequences (see, e.g., Karlin and Altschul (1993) Proc. Nat'l Acad. Sci. USA 90:5873-5787). One measure of similarity provided by the BLAST algorithm is the smallest sum probability (P(N)), which provides an indication of the probability by which a match between two nucleotide or amino acid sequences would occur by chance. For example, a nucleic acid is considered similar to a reference sequence if the smallest sum probability in a comparison of the test nucleic acid to the reference nucleic acid is less than about 0.1, more preferably less than about 0.01, and most preferably less than about 0.001.

A further indication that two nucleic acid sequences of polypeptides are substantially identical is that the polypeptide encoded by the first nucleic acid is immunologically cross reactive with the polypeptide encoded by the second nucleic acid, as described below. Thus, a polypeptide is typically substantially identical to a second polypeptide, e.g., where the two peptides differ only by conservative substitutions. Another indication that two nucleic acid sequences are substantially identical is that the two molecules hybridize to each other under stringent conditions, as described below.

The isolated DNA can be readily modified by nucleotide substitutions, nucleotide deletions, nucleotide insertions, and inversions of nucleotide stretches. These modifications result in novel DNA sequences which encode this protein or its derivatives. These modified sequences can be used to produce mutant proteins (muteins) or to enhance the expression of variant species. Enhanced

expression may involve gene amplification, increased transcription, increased translation, and other mechanisms. Such mutant DTLR-like derivatives include predetermined or site-specific mutations of the protein or its fragments, including silent mutations using genetic code degeneracy. "Mutant DTLR" as used herein encompasses a polypeptide otherwise falling within the homology definition of the DTLR as set forth above, but having an amino acid sequence which differs from that of other DTLR-like proteins as found in nature, whether by way of deletion, substitution, or insertion. In particular, "site specific mutant DTLR" encompasses a protein having substantial homology with a protein of Tables 2-10, and typically shares most of the biological activities or effects of the forms disclosed herein.

Although site specific mutation sites are predetermined, mutants need not be site specific. Mammalian DTLR mutagenesis can be achieved by making amino acid insertions or deletions in the gene, coupled with expression. Substitutions, deletions, insertions, or any combinations may be generated to arrive at a final construct. Insertions include amino- or carboxy- terminal fusions. Random mutagenesis can be conducted at a target codon and the expressed mammalian DTLR mutants can then be screened for the desired activity. Methods for making substitution mutations at predetermined sites in DNA having a known sequence are well known in the art, e.g., by M13 primer mutagenesis. See also Sambrook, et al. (1989) and Ausubel, et al. (1987 and periodic Supplements).

The mutations in the DNA normally should not place coding sequences out of reading frames and preferably will not create complementary regions that could hybridize to produce secondary mRNA structure such as loops or hairpins.

The phosphoramidite method described by Beaucage and Carruthers (1981) Tetra. Letts. 22:1859-1862, will produce suitable synthetic DNA fragments. A double stranded fragment will often be obtained either by synthesizing the complementary strand and annealing the strand together under appropriate conditions or by adding the complementary strand using DNA polymerase with an appropriate primer sequence.

Polymerase chain reaction (PCR) techniques can often be applied in mutagenesis. Alternatively, mutagenesis primers are commonly used methods for generating defined mutations at predetermined sites. See, e.g., Innis, et al. (eds. 1990) PCR Protocols: A Guide to Methods and Applications Academic Press, San Diego, CA; and Dieffenbach and Dveksler (eds. 1995) PCR Primer: A Laboratory Manual Cold Spring Harbor Press, CSH, NY.

IV. Proteins, Peptides

As described above, the present invention encompasses primate DTLR2-10, e.g., whose sequences are disclosed in Tables 2-10, and described above. Allelic and other variants are also contemplated, including, e.g., fusion proteins combining portions of such sequences with others, including epitope tags and functional domains.

The present invention also provides recombinant proteins, e.g., heterologous fusion proteins using segments from these rodent proteins. A heterologous fusion protein is a fusion of proteins or segments which are naturally not normally fused in the same manner. Thus, the fusion product of a DTLR with an IL-1 receptor is a continuous protein molecule having sequences fused in a typical peptide linkage, typically made as a single translation product and exhibiting properties, e.g., sequence or antigenicity, derived from each source peptide. A similar concept applies to heterologous nucleic acid sequences.

In addition, new constructs may be made from combining similar functional or structural domains from other related proteins, e.g., IL-1 receptors or other DTLRs, including species variants. For example, ligand-binding or other segments may be "swapped" between different new fusion polypeptides or fragments. See, e.g., Cunningham, et al. (1989) Science 243:1330-1336; and O'Dowd, et al. (1988) J. Biol. Chem. 263:15985-15992, each of which is incorporated herein by reference. Thus, new chimeric polypeptides exhibiting new combinations of specificities will result from the functional linkage of receptor-binding specificities. For example, the ligand binding domains from other related receptor molecules may be added or substituted for other domains of this or related proteins. The resulting protein will often have hybrid function and properties. For example, a fusion protein may include a targeting domain which may serve to provide sequestering of the fusion protein to a particular subcellular organelle.

Candidate fusion partners and sequences can be selected from various sequence data bases, e.g., GenBank, c/o IntelliGenetics, Mountain View, CA; and BCG, University of Wisconsin Biotechnology Computing Group, Madison, WI, which are each incorporated herein by reference.

The present invention particularly provides muteins which bind Toll ligands, and/or which are affected in signal transduction. Structural alignment of human DTLR1-10 with other members of the IL-1 family show conserved features/residues. See, e.g., Figure 3A. Alignment of the human DTLR sequences with other members of the IL-1 family indicates various structural and functionally shared features. See also, Bazan, et al. (1996) Nature 379:591; Lodi, et al. (1994) Science 263:1762-1766; Sayle and Milner-White (1995) TIBS 20:374-376; and Gronenberg, et al. (1991) Protein Engineering 4:263-269.

The IL-1α and IL-1β ligands bind an IL-1 receptor type I as the primary receptor and this complex then forms a high affinity receptor complex with the IL-1 receptor type III. Such receptor subunits are probably shared with the new IL-1 family members.

Similar variations in other species counterparts of DTLR2-10 sequences, e.g., in the corresponding regions, should provide similar interactions with ligand or substrate. Substitutions with either mouse sequences or human sequences are particularly preferred. Conversely, conservative substitutions away from the ligand binding interaction regions will probably preserve most signaling activities.

"Derivatives" of the primate DTLR2-10 include amino acid sequence mutants, glycosylation variants, metabolic derivatives and covalent or aggregative conjugates with other chemical moieties. Covalent derivatives can be prepared by linkage of functionalities to groups which are found in the DTLR amino acid side chains or at the N- or C- termini, e.g., by means which are well known in the art. These derivatives can include, without limitation, aliphatic esters or amides of the carboxyl terminus, or of residues containing carboxyl side chains, O-acyl derivatives of hydroxyl group-containing residues, and N-acyl derivatives of the amino terminal amino acid or amino-group containing residues, e.g., lysine or arginine. Acyl groups are selected from the group of alkyl-moieties including C3 to C18 normal alkyl, thereby forming alkanoyl aroyl species.

In particular, glycosylation alterations are included, e.g., made by modifying the glycosylation patterns of a polypeptide during its synthesis and processing, or in further processing steps. Particularly preferred means for accomplishing this are by exposing the polypeptide to glycosylating enzymes derived from cells which normally provide such processing, e.g., mammalian

glycosylation enzymes. Deglycosylation enzymes are also contemplated. Also embraced are versions of the same primary amino acid sequence which have other minor modifications, including phosphorylated amino acid residues, e.g., phosphotyrosine, phosphoserine, or phosphothreonine.

A major group of derivatives are covalent conjugates of the receptors or fragments thereof with other proteins of polypeptides. These derivatives can be synthesized in recombinant culture such as N- or C-terminal fusions or by the use of agents known in the art for their usefulness in cross-linking proteins through reactive side groups. Preferred derivatization sites with cross-linking agents are at free amino groups, carbohydrate moieties, and cysteine residues.

Fusion polypeptides between the receptors and other homologous or heterologous proteins are also provided. Homologous polypeptides may be fusions between different receptors, resulting in, for instance, a hybrid protein exhibiting binding specificity for multiple different Toll ligands, or a receptor which may have broadened or weakened specificity of substrate effect. Likewise, heterologous fusions may be constructed which would exhibit a combination of properties or activities of the derivative proteins. Typical examples are fusions of a reporter polypeptide, e.g., luciferase, with a segment or domain of a receptor, e.g., a ligand-binding segment, so that the presence or location of a desired ligand may be easily determined. See, e.g., Dull, et al., U.S. Patent No. 4,859,609, which is hereby incorporated herein by reference. Other gene fusion partners include glutathione-S-transferase (GST), bacterial ß-galactosidase, trpE, Protein A, ß-lactamase, alpha amylase, alcohol dehydrogenase, and yeast alpha mating factor. See, e.g., Godowski, et al. (1988) Science 241:812-816.

The phosphoramidite method described by Beaucage and Carruthers (1981) Tetra. Letts. 22:1859-1862, will produce suitable synthetic DNA fragments. A double stranded fragment will often be obtained either by synthesizing the complementary strand and annealing the strand together under appropriate conditions or by adding the complementary strand using DNA polymerase with an appropriate primer sequence.

Such polypeptides may also have amino acid residues which have been chemically modified by phosphorylation, sulfonation, biotinylation, or the addition or removal of other moieties, particularly those which have molecular shapes similar to phosphate groups. In some embodiments, the modifications will be useful labeling reagents, or serve as purification targets, e.g., affinity ligands.

Fusion proteins will typically be made by either recombinant nucleic acid methods or by synthetic polypeptide methods. Techniques for nucleic acid manipulation and expression are described generally, for example, in Sambrook, et al. (1989) Molecular Cloning: A Laboratory Manual (2d ed.), Vols. 1-3, Cold Spring Harbor Laboratory, and Ausubel, et al. (eds. 1987 and periodic supplements) Current Protocols in Molecular Biology, Greene/Wiley, New York, which are each incorporated herein by reference. Techniques for synthesis of polypeptides are described, for example, in Merrifield (1963) J. Amer. Chem. Soc. 85:2149-2156; Merrifield (1986) Science 232: 341-347; and Atherton, et al. (1989) Solid Phase Peptide Synthesis: A Practical Approach, IRL Press, Oxford; each of which is incorporated herein by reference. See also Dawson, et al. (1994) Science 266:776-779 for methods to make larger polypeptides.

This invention also contemplates the use of derivatives of a DTLR2-10 other than variations in amino acid sequence or glycosylation. Such derivatives may involve covalent or aggregative association with chemical

moieties. These derivatives generally fall into three classes: (1) salts, (2) side chain and terminal residue covalent modifications, and (3) adsorption complexes, for example with cell membranes. Such covalent or aggregative derivatives are useful as immunogens, as reagents in immunoassays, or in purification methods such as for affinity purification of a receptor or other binding molecule, e.g., an antibody. For example, a Toll ligand can be immobilized by covalent bonding to a solid support such as cyanogen bromide-activated Sepharose, by methods which are well known in the art, or adsorbed onto polyolefin surfaces, with or without glutaraldehyde cross-linking, for use in the assay or purification of a DTLR receptor, antibodies, or other similar molecules. The ligand can also be labeled with a detectable group, for example radioiodinated by the chloramine T procedure, covalently bound to rare earth chelates, or conjugated to another fluorescent moiety for use in diagnostic assays.

A DTLR of this invention can be used as an immunogen for the production of antisera or antibodies specific, e.g., capable of distinguishing between other IL-1 receptor family members, for the DTLR or various fragments thereof. The purified DTLR can be used to screen monoclonal antibodies or antigen-binding fragments prepared by immunization with various forms of impure preparations containing the protein. In particular, the term "antibodies" also encompasses antigen binding fragments of natural antibodies, e.g., Fab, Fab2, Fv, etc. The purified DTLR can also be used as a reagent to detect antibodies generated in response to the presence of elevated levels of expression, or immunological disorders which lead to antibody production to the endogenous receptor. Additionally, DTLR fragments may also serve as immunogens to produce the antibodies of the present invention, as described immediately below. For example, this invention contemplates antibodies having binding

affinity to or being raised against the amino acid sequences shown in Tables 2-10, fragments thereof, or various homologous peptides. In particular, this invention contemplates antibodies having binding affinity to, or having been raised against, specific fragments which are predicted to be, or actually are, exposed at the exterior protein surface of the native DTLR.

The blocking of physiological response to the receptor ligands may result from the inhibition of binding of the ligand to the receptor, likely through competitive inhibition. Thus, in vitro assays of the present invention will often use antibodies or antigen binding segments of these antibodies, or fragments attached to solid phase substrates. These assays will also allow for the diagnostic determination of the effects of either ligand binding region mutations and modifications, or other mutations and modifications, e.g., which affect signaling or enzymatic function.

This invention also contemplates the use of competitive drug screening assays, e.g., where neutralizing antibodies to the receptor or fragments compete with a test compound for binding to a ligand or other antibody. In this manner, the neutralizing antibodies or fragments can be used to detect the presence of a polypeptide which shares one or more binding sites to a receptor and can also be used to occupy binding sites on a receptor that might otherwise bind a ligand.

V. Making Nucleic Acids and Protein

DNA which encodes the protein or fragments thereof can be obtained by chemical synthesis, screening cDNA libraries, or by screening genomic libraries prepared from a wide variety of cell lines or tissue samples. Natural sequences can be isolated using standard methods and the sequences provided herein, e.g., in Tables 2-10. Other species counterparts can be identified by hybridization

techniques, or by various PCR techniques, combined with or by searching in sequence databases, e.g., GenBank.

This DNA can be expressed in a wide variety of host cells for the synthesis of a full-length receptor or fragments which can in turn, for example, be used to generate polyclonal or monoclonal antibodies; for binding studies; for construction and expression of modified ligand binding or kinase/phosphatase domains; and for structure/function studies. Variants or fragments can be expressed in host cells that are transformed or transfected with appropriate expression vectors. These molecules can be substantially free of protein or cellular contaminants, other than those derived from the recombinant host, and therefore are particularly useful in pharmaceutical compositions when combined with a pharmaceutically acceptable carrier and/or diluent. The protein, or portions thereof, may be expressed as fusions with other proteins.

Expression vectors are typically self-replicating DNA or RNA constructs containing the desired receptor gene or its fragments, usually operably linked to suitable genetic control elements that are recognized in a suitable host cell. These control elements are capable of effecting expression within a suitable host. The specific type of control elements necessary to effect expression will depend upon the eventual host cell used. Generally, the genetic control elements can include a prokaryotic promoter system or a eukaryotic promoter expression control system, and typically include a transcriptional promoter, an optional operator to control the onset of transcription, transcription enhancers to elevate the level of mRNA expression, a sequence that encodes a suitable ribosome binding site, and sequences that terminate transcription and translation. Expression vectors also usually contain an origin of replication that

allows the vector to replicate independently of the host cell.

The vectors of this invention include those which contain DNA which encodes a protein, as described, or a fragment thereof encoding a biologically active equivalent polypeptide. The DNA can be under the control of a viral promoter and can encode a selection marker. This invention further contemplates use of such expression vectors which are capable of expressing eukaryotic cDNA coding for such a protein in a prokaryotic or eukaryotic host, where the vector is compatible with the host and where the eukaryotic cDNA coding for the receptor is inserted into the vector such that growth of the host containing the vector expresses the cDNA in question. Usually, expression vectors are designed for stable replication in their host cells or for amplification to greatly increase the total number of copies of the desirable gene per cell. It is not always necessary to require that an expression vector replicate in a host cell, e.g., it is possible to effect transient expression of the protein or its fragments in various hosts using vectors that do not contain a replication origin that is recognized by the host cell. It is also possible to use vectors that cause integration of the protein encoding portion or its fragments into the host DNA by recombination.

Vectors, as used herein, comprise plasmids, viruses, bacteriophage, integratable DNA fragments, and other vehicles which enable the integration of DNA fragments into the genome of the host. Expression vectors are specialized vectors which contain genetic control elements that effect expression of operably linked genes. Plasmids are the most commonly used form of vector but all other forms of vectors which serve an equivalent function and which are, or become, known in the art are suitable for use herein. See, e.g., Pouwels, et al. (1985 and

Supplements) Cloning Vectors: A Laboratory Manual, Elsevier, N.Y., and Rodriquez, et al. (eds.) Vectors: A Survey of Molecular Cloning Vectors and Their Uses, Buttersworth, Boston, 1988, which are incorporated herein by reference.

Transformed cells are cells, preferably mammalian, that have been transformed or transfected with receptor vectors constructed using recombinant DNA techniques. Transformed host cells usually express the desired protein or its fragments, but for purposes of cloning, amplifying, and manipulating its DNA, do not need to express the subject protein. This invention further contemplates culturing transformed cells in a nutrient medium, thus permitting the receptor to accumulate in the cell membrane. The protein can be recovered, either from the culture or, in certain instances, from the culture medium.

For purposes of this invention, nucleic sequences are operably linked when they are functionally related to each other. For example, DNA for a presequence or secretory leader is operably linked to a polypeptide if it is expressed as a preprotein or participates in directing the polypeptide to the cell membrane or in secretion of the polypeptide. A promoter is operably linked to a coding sequence if it controls the transcription of the polypeptide; a ribosome binding site is operably linked to a coding sequence if it is positioned to permit translation. Usually, operably linked means contiguous and in reading frame, however, certain genetic elements such as repressor genes are not contiguously linked but still bind to operator sequences that in turn control expression.

Suitable host cells include prokaryotes, lower eukaryotes, and higher eukaryotes. Prokaryotes include both gram negative and gram positive organisms, e.g., E. coli and B. subtilis. Lower eukaryotes include yeasts, e.g., S. cerevisiae and Pichia, and species of the genus

Dictyostelium. Higher eukaryotes include established tissue culture cell lines from animal cells, both of non-mammalian origin, e.g., insect cells, and birds, and of mammalian origin, e.g., human, primates, and rodents.

Prokaryotic host-vector systems include a wide variety of vectors for many different species. As used herein, E. coli and its vectors will be used generically to include equivalent vectors used in other prokaryotes. A representative vector for amplifying DNA is pBR322 or many of its derivatives. Vectors that can be used to express the receptor or its fragments include, but are not limited to, such vectors as those containing the lac promoter (pUC-series); trp promoter (pBR322-trp); Ipp promoter (the pIN-series); lambda-pP or pR promoters (pOTS); or hybrid promoters such as ptac (pDR540). See Brosius, et al. (1988) "Expression Vectors Employing Lambda-, trp-, lac-, and Ipp-derived Promoters", in Vectors: A Survey of Molecular Cloning Vectors and Their Uses, (eds. Rodriguez and Denhardt), Buttersworth, Boston, Chapter 10, pp. 205-236, which is incorporated herein by reference.

Lower eukaryotes, e.g., yeasts and Dictyostelium, may be transformed with DTLR sequence containing vectors. For purposes of this invention, the most common lower eukaryotic host is the baker's yeast, Saccharomyces cerevisiae. It will be used to generically represent lower eukaryotes although a number of other strains and species are also available. Yeast vectors typically consist of a replication origin (unless of the integrating type), a selection gene, a promoter, DNA encoding the receptor or its fragments, and sequences for translation termination, polyadenylation, and transcription termination. Suitable expression vectors for yeast include such constitutive promoters as 3-phosphoglycerate kinase and various other glycolytic enzyme gene promoters or such inducible promoters as the alcohol dehydrogenase 2

promoter or metallothionine promoter. Suitable vectors include derivatives of the following types: self-replicating low copy number (such as the YRp-series), self-replicating high copy number (such as the YEp-series); integrating types (such as the YIp-series), or mini-chromosomes (such as the YCp-series).

Higher eukaryotic tissue culture cells are normally the preferred host cells for expression of the functionally active interleukin protein. In principle, any higher eukaryotic tissue culture cell line is workable, e.g., insect baculovirus expression systems, whether from an invertebrate or vertebrate source. However, mammalian cells are preferred. Transformation or transfection and propagation of such cells has become a routine procedure. Examples of useful cell lines include HeLa cells, Chinese hamster ovary (CHO) cell lines, baby rat kidney (BRK) cell lines, insect cell lines, bird cell lines, and monkey (COS) cell lines. Expression vectors for such cell lines usually include an origin of replication, a promoter, a translation initiation site, RNA splice sites (if genomic DNA is used), a polyadenylation site, and a transcription termination site. These vectors also usually contain a selection gene or amplification gene. Suitable expression vectors may be plasmids, viruses, or retroviruses carrying promoters derived, e.g., from such sources as from adenovirus, SV40, parvoviruses, vaccinia virus, or cytomegalovirus. Representative examples of suitable expression vectors include pCDNA1; pCD, see Okayama, et al. (1985) Mol. Cell Biol. 5:1136-1142; pMC1neo PolyA, see Thomas, et al. (1987) Cell 51:503-512; and a baculovirus vector such as pAC 373 or pAC 610.

For secreted proteins, an open reading frame usually encodes a polypeptide that consists of a mature or secreted product covalently linked at its N-terminus to a signal peptide. The signal peptide is cleaved prior to

secretion of the mature, or active, polypeptide. The cleavage site can be predicted with a high degree of accuracy from empirical rules, e.g., von-Heijne (1986) Nucleic Acids Research 14:4683-4690, and the precise amino acid composition of the signal peptide does not appear to be critical to its function, e.g., Randall, et al. (1989) Science 243:1156-1159; Kaiser, et al. (1987) Science 235:312-317.

It will often be desired to express these polypeptides in a system which provides a specific or defined glycosylation pattern. In this case, the usual pattern will be that provided naturally by the expression system. However, the pattern will be modifiable by exposing the polypeptide, e.g., an unglycosylated form, to appropriate glycosylating proteins introduced into a heterologous expression system. For example, the receptor gene may be co-transformed with one or more genes encoding mammalian or other glycosylating enzymes. Using this approach, certain mammalian glycosylation patterns will be achievable in prokaryote or other cells.

The source of DTLR can be a eukaryotic or prokaryotic host expressing recombinant DTLR, such as is described above. The source can also be a cell line such as mouse Swiss 3T3 fibroblasts, but other mammalian cell lines are also contemplated by this invention, with the preferred cell line being from the human species.

Now that the sequences are known, the primate DTLRs, fragments, or derivatives thereof can be prepared by conventional processes for synthesizing peptides. These include processes such as are described in Stewart and Young (1984) Solid Phase Peptide Synthesis, Pierce Chemical Co., Rockford, IL; Bodanszky and Bodanszky (1984) The Practice of Peptide Synthesis, Springer-Verlag, New York; and Bodanszky (1984) The Principles of Peptide Synthesis, Springer-Verlag, New York; all of each which are incorporated herein by reference. For example, an

azide process, an acid chloride process, an acid anhydride process, a mixed anhydride process, an active ester process (e.g., p-nitrophenyl ester, N-hydroxysuccinimide ester, or cyanomethyl ester), a carbodiimidazole process, an oxidative-reductive process, or a dicyclohexylcarbodiimide (DCCD)/additive process can be used. Solid phase and solution phase syntheses are both applicable to the foregoing processes. Similar techniques can be used with partial DTLR sequences.

The DTLR proteins, fragments, or derivatives are suitably prepared in accordance with the above processes as typically employed in peptide synthesis, generally either by a so-called stepwise process which comprises condensing an amino acid to the terminal amino acid, one by one in sequence, or by coupling peptide fragments to the terminal amino acid. Amino groups that are not being used in the coupling reaction typically must be protected to prevent coupling at an incorrect location.

If a solid phase synthesis is adopted, the C-terminal amino acid is bound to an insoluble carrier or support through its carboxyl group. The insoluble carrier is not particularly limited as long as it has a binding capability to a reactive carboxyl group. Examples of such insoluble carriers include halomethyl resins, such as chloromethyl resin or bromomethyl resin, hydroxymethyl resins, phenol resins, tert-alkyloxycarbonylhydrazidated resins, and the like.

An amino group-protected amino acid is bound in sequence through condensation of its activated carboxyl group and the reactive amino group of the previously formed peptide or chain, to synthesize the peptide step by step. After synthesizing the complete sequence, the peptide is split off from the insoluble carrier to produce the peptide. This solid-phase approach is generally described by Merrifield, et al. (1963) in J. Am. Chem.

Soc. 85:2149-2156, which is incorporated herein by reference.

The prepared protein and fragments thereof can be isolated and purified from the reaction mixture by means of peptide separation, for example, by extraction, precipitation, electrophoresis, various forms of chromatography, and the like. The receptors of this invention can be obtained in varying degrees of purity depending upon desired uses. Purification can be accomplished by use of the protein purification techniques disclosed herein, see below, or by the use of the antibodies herein described in methods of immunoabsorbant affinity chromatography. This immunoabsorbant affinity chromatography is carried out by first linking the antibodies to a solid support and then contacting the linked antibodies with solubilized lysates of appropriate cells, lysates of other cells expressing the receptor, or lysates or supernatants of cells producing the protein as a result of DNA techniques, see below.

Generally, the purified protein will be at least about 40% pure, ordinarily at least about 50% pure, usually at least about 60% pure, typically at least about 70% pure, more typically at least about 80% pure, preferable at least about 90% pure and more preferably at least about 95% pure, and in particular embodiments, 97%-99% or more. Purity will usually be on a weight basis, but can also be on a molar basis. Different assays will be applied as appropriate.

VI. Antibodies

Antibodies can be raised to the various mammalian, e.g., primate DTLR proteins and fragments thereof, both in naturally occurring native forms and in their recombinant forms, the difference being that antibodies to the active receptor are more likely to recognize epitopes which are only present in the native conformations. Denatured

antigen detection can also be useful in, e.g., Western analysis. Anti-idiotypic antibodies are also contemplated, which would be useful as agonists or antagonists of a natural receptor or an antibody.

Preferred antibodies will exhibit properties of both affinity and selectivity. High affinity is generally preferred, while selectivity will allow distinction between various embodiment subsets. In particular, it will be desirable to possess antibody preparations characterized to bind, e.g., various specific combinations of related members while not binding others. Such various combinatorial subsets are specifically enabled, e.g., these reagents may be generated or selected using standard methods of immunoaffinity, selection, etc.

Antibodies, including binding fragments and single chain versions, against predetermined fragments of the protein can be raised by immunization of animals with conjugates of the fragments with immunogenic proteins. Monoclonal antibodies are prepared from cells secreting the desired antibody. These antibodies can be screened for binding to normal or defective protein, or screened for agonistic or antagonistic activity. These monoclonal antibodies will usually bind with at least a K_D of about 1 mM, more usually at least about 300 µM, typically at least about 100µM, more typically at least about 30 µM, preferably at least about 10 µM, and more preferably at least about 3 µM or better.

The antibodies, including antigen binding fragments, of this invention can have significant diagnostic or therapeutic value. They can be potent antagonists that bind to the receptor and inhibit binding to ligand or inhibit the ability of the receptor to elicit a biological response, e.g., act on its substrate. They also can be useful as non-neutralizing antibodies and can be coupled to toxins or radionuclides to bind producing cells, or cells localized to the source of the interleukin.

Further, these antibodies can be conjugated to drugs or other therapeutic agents, either directly or indirectly by means of a linker.

The antibodies of this invention can also be useful in diagnostic applications. As capture or non-neutralizing antibodies, they might bind to the receptor without inhibiting ligand or substrate binding. As neutralizing antibodies, they can be useful in competitive binding assays. They will also be useful in detecting or quantifying ligand. They may be used as reagents for Western blot analysis, or for immunoprecipitation or immunopurification of the respective protein.

Protein fragments may be joined to other materials, particularly polypeptides, as fused or covalently joined polypeptides to be used as immunogens. Mammalian DTLR and its fragments may be fused or covalently linked to a variety of immunogens, such as keyhole limpet hemocyanin, bovine serum albumin, tetanus toxoid, etc. See Microbiology, Hoeber Medical Division, Harper and Row, 1969; Landsteiner (1962) Specificity of Serological Reactions, Dover Publications, New York; and Williams, et al. (1967) Methods in Immunology and Immunochemistry, Vol. 1, Academic Press, New York; each of which are incorporated herein by reference, for descriptions of methods of preparing polyclonal antisera. A typical method involves hyperimmunization of an animal with an antigen. The blood of the animal is then collected shortly after the repeated immunizations and the gamma globulin is isolated.

In some instances, it is desirable to prepare monoclonal antibodies from various mammalian hosts, such as mice, rodents, primates, humans, etc. Description of techniques for preparing such monoclonal antibodies may be found in, e.g., Stites, et al. (eds.) Basic and Clinical Immunology (4th ed.), Lange Medical Publications, Los

Altos, CA, and references cited therein; Harlow and Lane (1988) Antibodies: A Laboratory Manual, CSH Press; Goding (1986) Monoclonal Antibodies: Principles and Practice (2d ed) Academic Press, New York; and particularly in Kohler and Milstein (1975) in Nature 256: 495-497, which discusses one method of generating monoclonal antibodies. Each of these references is incorporated herein by reference. Summarized briefly, this method involves injecting an animal with an immunogen. The animal is then sacrificed and cells taken from its spleen, which are then fused with myeloma cells. The result is a hybrid cell or "hybridoma" that is capable of reproducing in vitro. The population of hybridomas is then screened to isolate individual clones, each of which secrete a single antibody species to the immunogen. In this manner, the individual antibody species obtained are the products of immortalized and cloned single B cells from the immune animal generated in response to a specific site recognized on the immunogenic substance.

Other suitable techniques involve in vitro exposure of lymphocytes to the antigenic polypeptides or alternatively to selection of libraries of antibodies in phage or similar vectors. See, Huse, et al. (1989) "Generation of a Large Combinatorial Library of the Immunoglobulin Repertoire in Phage Lambda," Science 246:1275-1281; and Ward, et al. (1989) Nature 341:544-546, each of which is hereby incorporated herein by reference. The polypeptides and antibodies of the present invention may be used with or without modification, including chimeric or humanized antibodies. Frequently, the polypeptides and antibodies will be labeled by joining, either covalently or non-covalently, a substance which provides for a detectable signal. A wide variety of labels and conjugation techniques are known and are reported extensively in both the scientific and patent literature. Suitable labels include radionuclides,

enzymes, substrates, cofactors, inhibitors, fluorescent moieties, chemiluminescent moieties, magnetic particles, and the like. Patents, teaching the use of such labels include U.S. Patent Nos. 3,817,837; 3,850,752; 3,939,350; 3,996,345; 4,277,437; 4,275,149; and 4,366,241. Also, recombinant or chimeric immunoglobulins may be produced, see Cabilly, U.S. Patent No. 4,816,567; or made in transgenic mice, see Mendez, et al. (1997) Nature Genetics 15:146-156. These references are incorporated herein by reference.

The antibodies of this invention can also be used for affinity chromatography in isolating the DTLRs. Columns can be prepared where the antibodies are linked to a solid support, e.g., particles, such as agarose, Sephadex, or the like, where a cell lysate may be passed through the column, the column washed, followed by increasing concentrations of a mild denaturant, whereby the purified protein will be released. The protein may be used to purify antibody.

The antibodies may also be used to screen expression libraries for particular expression products. Usually the antibodies used in such a procedure will be labeled with a moiety allowing easy detection of presence of antigen by antibody binding.

Antibodies raised against a DTLR will also be used to raise anti-idiotypic antibodies. These will be useful in detecting or diagnosing various immunological conditions related to expression of the protein or cells which express the protein. They also will be useful as agonists or antagonists of the ligand, which may be competitive inhibitors or substitutes for naturally occurring ligands.

A DTLR protein that specifically binds to or that is specifically immunoreactive with an antibody generated against a defined immunogen, such as an immunogen consisting of the amino acid sequence of SEQ ID NO: 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, or 24, is typically

determined in an immunoassay. The immunoassay typically uses a polyclonal antiserum which was raised, e.g., to a protein of SEQ ID NO: 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, or 24. This antiserum is selected to have low crossreactivity against other IL-1R family members, e.g., DTLR1, preferably from the same species, and any such crossreactivity is removed by immunoabsorption prior to use in the immunoassay.

In order to produce antisera for use in an immunoassay, the protein of SEQ ID NO: 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, or 24, or a combination thereof, is isolated as described herein. For example, recombinant protein may be produced in a mammalian cell line. An appropriate host, e.g., an inbred strain of mice such as Balb/c, is immunized with the selected protein, typically using a standard adjuvant, such as Freund's adjuvant, and a standard mouse immunization protocol (see Harlow and Lane, supra). Alternatively, a synthetic peptide derived from the sequences disclosed herein and conjugated to a carrier protein can be used an immunogen. Polyclonal sera are collected and titered against the immunogen protein in an immunoassay, e.g., a solid phase immunoassay with the immunogen immobilized on a solid support. Polyclonal antisera with a titer of 10^4 or greater are selected and tested for their cross reactivity against other IL-1R family members, e.g., mouse DTLRs or human DTLR1, using a competitive binding immunoassay such as the one described in Harlow and Lane, supra, at pages 570-573. Preferably at least two DTLR family members are used in this determination in conjunction with either or some of the human DTLR2-10. These IL-1R family members can be produced as recombinant proteins and isolated using standard molecular biology and protein chemistry techniques as described herein.

Immunoassays in the competitive binding format can be used for the crossreactivity determinations. For example,

the proteins of SEQ ID NO: 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, and/or 24, or various fragments thereof, can be immobilized to a solid support. Proteins added to the assay compete with the binding of the antisera to the immobilized antigen. The ability of the above proteins to compete with the binding of the antisera to the immobilized protein is compared to the protein of SEQ ID NO: 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, and/or 24. The percent crossreactivity for the above proteins is calculated, using standard calculations. Those antisera with less than 10% crossreactivity with each of the proteins listed above are selected and pooled. The cross-reacting antibodies are then removed from the pooled antisera by immunoabsorption with the above-listed proteins.

The immunoabsorbed and pooled antisera are then used in a competitive binding immunoassay as described above to compare a second protein to the immunogen protein (e.g., the IL-1R like protein of SEQ ID NO: 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, and/or 24). In order to make this comparison, the two proteins are each assayed at a wide range of concentrations and the amount of each protein required to inhibit 50% of the binding of the antisera to the immobilized protein is determined. If the amount of the second protein required is less than twice the amount of the protein of the selected protein or proteins that is required, then the second protein is said to specifically bind to an antibody generated to the immunogen.

It is understood that these DTLR proteins are members of a family of homologous proteins that comprise at least 10 so far identified genes. For a particular gene product, such as the DTLR2-10, the term refers not only to the amino acid sequences disclosed herein, but also to other proteins that are allelic, non-allelic or species variants. It also understood that the terms include nonnatural mutations introduced by deliberate mutation

using conventional recombinant technology such as single site mutation, or by excising short sections of DNA encoding the respective proteins, or by substituting new amino acids, or adding new amino acids. Such minor alterations must substantially maintain the immunoidentity of the original molecule and/or its biological activity. Thus, these alterations include proteins that are specifically immunoreactive with a designated naturally occurring IL-1R related protein, for example, the DTLR proteins shown in SEQ ID NO: 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, or 24. The biological properties of the altered proteins can be determined by expressing the protein in an appropriate cell line and measuring the appropriate effect upon lymphocytes. Particular protein modifications considered minor would include conservative substitution of amino acids with similar chemical properties, as described above for the IL-1R family as a whole. By aligning a protein optimally with the protein of DTLR2-10 and by using the conventional immunoassays described herein to determine immunoidentity, one can determine the protein compositions of the invention.

VII. Kits and quantitation

Both naturally occurring and recombinant forms of the IL-1R like molecules of this invention are particularly useful in kits and assay methods. For example, these methods would also be applied to screening for binding activity, e.g., ligands for these proteins. Several methods of automating assays have been developed in recent years so as to permit screening of tens of thousands of compounds per year. See, e.g., a BIOMEK automated workstation, Beckman Instruments, Palo Alto, California, and Fodor, et al. (1991) Science 251:767-773, which is incorporated herein by reference. The latter describes means for testing binding by a plurality of defined polymers synthesized on a solid substrate. The

development of suitable assays to screen for a ligand or agonist/antagonist homologous proteins can be greatly facilitated by the availability of large amounts of purified, soluble DTLRs in an active state such as is provided by this invention.

Purified DTLR can be coated directly onto plates for use in the aforementioned ligand screening techniques. However, non-neutralizing antibodies to these proteins can be used as capture antibodies to immobilize the respective receptor on the solid phase, useful, e.g., in diagnostic uses.

This invention also contemplates use of DTLR2-10, fragments thereof, peptides, and their fusion products in a variety of diagnostic kits and methods for detecting the presence of the protein or its ligand. Alternatively, or additionally, antibodies against the molecules may be incorporated into the kits and methods. Typically the kit will have a compartment containing either a defined DTLR peptide or gene segment or a reagent which recognizes one or the other. Typically, recognition reagents, in the case of peptide, would be a receptor or antibody, or in the case of a gene segment, would usually be a hybridization probe.

A preferred kit for determining the concentration of, e.g., DTLR4, a sample would typically comprise a labeled compound, e.g., ligand or antibody, having known binding affinity for DTLR4, a source of DTLR4 (naturally occurring or recombinant) as a positive control, and a means for separating the bound from free labeled compound, for example a solid phase for immobilizing the DTLR4 in the test sample. Compartments containing reagents, and instructions, will normally be provided.

Antibodies, including antigen binding fragments, specific for mammalian DTLR or a peptide fragment, or receptor fragments are useful in diagnostic applications to detect the presence of elevated levels of ligand and/or

its fragments. Diagnostic assays may be homogeneous (without a separation step between free reagent and antibody-antigen complex) or heterogeneous (with a separation step). Various commercial assays exist, such as radioimmunoassay (RIA), enzyme-linked immunosorbent assay (ELISA), enzyme immunoassay (EIA), enzyme-multiplied immunoassay technique (EMIT), substrate-labeled fluorescent immunoassay (SLFIA) and the like. For example, unlabeled antibodies can be employed by using a second antibody which is labeled and which recognizes the antibody to DTLR4 or to a particular fragment thereof. These assays have also been extensively discussed in the literature. See, e.g., Harlow and Lane (1988) Antibodies: A Laboratory Manual, CSH., and Coligan (ed. 1991 and periodic supplements) Current Protocols In Immunology Greene/Wiley, New York.

Anti-idiotypic antibodies may have similar use to serve as agonists or antagonists of DTLR4. These should be useful as therapeutic reagents under appropriate circumstances.

Frequently, the reagents for diagnostic assays are supplied in kits, so as to optimize the sensitivity of the assay. For the subject invention, depending upon the nature of the assay, the protocol, and the label, either labeled or unlabeled antibody, or labeled ligand is provided. This is usually in conjunction with other additives, such as buffers, stabilizers, materials necessary for signal production such as substrates for enzymes, and the like. Preferably, the kit will also contain instructions for proper use and disposal of the contents after use. Typically the kit has compartments for each useful reagent, and will contain instructions for proper use and disposal of reagents. Desirably, the reagents are provided as a dry lyophilized powder, where the reagents may be reconstituted in an aqueous medium

having appropriate concentrations for performing the assay.

The aforementioned constituents of the diagnostic assays may be used without modification or may be modified in a variety of ways. For example, labeling may be achieved by covalently or non-covalently joining a moiety which directly or indirectly provides a detectable signal. In any of these assays, a test compound, DTLR, or antibodies thereto can be labeled either directly or indirectly. Possibilities for direct labeling include label groups: radiolabels such as ^{125}I, enzymes (U.S. Pat. No. 3,645,090) such as peroxidase and alkaline phosphatase, and fluorescent labels (U.S. Pat. No. 3,940,475) capable of monitoring the change in fluorescence intensity, wavelength shift, or fluorescence polarization. Both of the patents are incorporated herein by reference. Possibilities for indirect labeling include biotinylation of one constituent followed by binding to avidin coupled to one of the above label groups.

There are also numerous methods of separating the bound from the free ligand, or alternatively the bound from the free test compound. The DTLR can be immobilized on various matrixes followed by washing. Suitable matrices include plastic such as an ELISA plate, filters, and beads. Methods of immobilizing the receptor to a matrix include, without limitation, direct adhesion to plastic, use of a capture antibody, chemical coupling, and biotin-avidin. The last step in this approach involves the precipitation of antibody/antigen complex by any of several methods including those utilizing, e.g., an organic solvent such as polyethylene glycol or a salt such as ammonium sulfate. Other suitable separation techniques include, without limitation, the fluorescein antibody magnetizable particle method described in Rattle, et al. (1984) Clin. Chem. 30(9):1457-1461, and the double antibody magnetic particle separation as described in U.S.

Pat. No. 4,659,678, each of which is incorporated herein by reference.

The methods for linking protein or fragments to various labels have been extensively reported in the literature and do not require detailed discussion here. Many of the techniques involve the use of activated carboxyl groups either through the use of carbodiimide or active esters to form peptide bonds, the formation of thioethers by reaction of a mercapto group with an activated halogen such as chloroacetyl, or an activated olefin such as maleimide, for linkage, or the like. Fusion proteins will also find use in these applications.

Another diagnostic aspect of this invention involves use of oligonucleotide or polynucleotide sequences taken from the sequence of a DTLR. These sequences can be used as probes for detecting levels of the respective DTLR in patients suspected of having an immunological disorder. The preparation of both RNA and DNA nucleotide sequences, the labeling of the sequences, and the preferred size of the sequences has received ample description and discussion in the literature. Normally an oligonucleotide probe should have at least about 14 nucleotides, usually at least about 18 nucleotides, and the polynucleotide probes may be up to several kilobases. Various labels may be employed, most commonly radionuclides, particularly ^{32}P. However, other techniques may also be employed, such as using biotin modified nucleotides for introduction into a polynucleotide. The biotin then serves as the site for binding to avidin or antibodies, which may be labeled with a wide variety of labels, such as radionuclides, fluorescers, enzymes, or the like. Alternatively, antibodies may be employed which can recognize specific duplexes, including DNA duplexes, RNA duplexes, DNA-RNA hybrid duplexes, or DNA-protein duplexes. The antibodies in turn may be labeled and the assay carried out where the duplex is bound to a surface, so that upon the formation

of duplex on the surface, the presence of antibody bound to the duplex can be detected. The use of probes to the novel anti-sense RNA may be carried out in any conventional techniques such as nucleic acid hybridization, plus and minus screening, recombinational probing, hybrid released translation (HRT), and hybrid arrested translation (HART). This also includes amplification techniques such as polymerase chain reaction (PCR).

Diagnostic kits which also test for the qualitative or quantitative presence of other markers are also contemplated. Diagnosis or prognosis may depend on the combination of multiple indications used as markers. Thus, kits may test for combinations of markers. See, e.g., Viallet, et al. (1989) Progress in Growth Factor Res. 1:89-97.

VIII. Therapeutic Utility

This invention provides reagents with significant therapeutic value. The DTLRs (naturally occurring or recombinant), fragments thereof, mutein receptors, and antibodies, along with compounds identified as having binding affinity to the receptors or antibodies, should be useful in the treatment of conditions exhibiting abnormal expression of the receptors of their ligands. Such abnormality will typically be manifested by immunological disorders. Additionally, this invention should provide therapeutic value in various diseases or disorders associated with abnormal expression or abnormal triggering of response to the ligand. The Toll ligands have been suggested to be involved in morphologic development, e.g., dorso-ventral polarity determination, and immune responses, particularly the primitive innate responses. See, e.g., Sun, et al. (1991) Eur. J. Biochem. 196:247-254; Hultmark (1994) Nature 367:116-117.

Recombinant DTLRs, muteins, agonist or antagonist antibodies thereto, or antibodies can be purified and then administered to a patient. These reagents can be combined for therapeutic use with additional active ingredients, e.g., in conventional pharmaceutically acceptable carriers or diluents, along with physiologically innocuous stabilizers and excipients. These combinations can be sterile, e.g., filtered, and placed into dosage forms as by lyophilization in dosage vials or storage in stabilized aqueous preparations. This invention also contemplates use of antibodies or binding fragments thereof which are not complement binding.

Ligand screening using DTLR or fragments thereof can be performed to identify molecules having binding affinity to the receptors. Subsequent biological assays can then be utilized to determine if a putative ligand can provide competitive binding, which can block intrinsic stimulating activity. Receptor fragments can be used as a blocker or antagonist in that it blocks the activity of ligand. Likewise, a compound having intrinsic stimulating activity can activate the receptor and is thus an agonist in that it simulates the activity of ligand, e.g., inducing signaling. This invention further contemplates the therapeutic use of antibodies to DTLRs as antagonists.

The quantities of reagents necessary for effective therapy will depend upon many different factors, including means of administration, target site, physiological state of the patient, and other medicants administered. Thus, treatment dosages should be titrated to optimize safety and efficacy. Typically, dosages used in vitro may provide useful guidance in the amounts useful for in situ administration of these reagents. Animal testing of effective doses for treatment of particular disorders will provide further predictive indication of human dosage. Various considerations are described, e.g., in Gilman, et al. (eds. 1990) Goodman and Gilman's: The Pharmacological

Bases of Therapeutics, 8th Ed., Pergamon Press; and Remington's Pharmaceutical Sciences, (current edition), Mack Publishing Co., Easton, Penn.; each of which is hereby incorporated herein by reference. Methods for administration are discussed therein and below, e.g., for oral, intravenous, intraperitoneal, or intramuscular administration, transdermal diffusion, and others. Pharmaceutically acceptable carriers will include water, saline, buffers, and other compounds described, e.g., in the Merck Index, Merck & Co., Rahway, New Jersey. Because of the likely high affinity binding, or turnover numbers, between a putative ligand and its receptors, low dosages of these reagents would be initially expected to be effective. And the signaling pathway suggests extremely low amounts of ligand may have effect. Thus, dosage ranges would ordinarily be expected to be in amounts lower than 1 mM concentrations, typically less than about 10 µM concentrations, usually less than about 100 nM, preferably less than about 10 pM (picomolar), and most preferably less than about 1 fM (femtomolar), with an appropriate carrier. Slow release formulations, or slow release apparatus will often be utilized for continuous administration.

DTLRs, fragments thereof, and antibodies or its fragments, antagonists, and agonists, may be administered directly to the host to be treated or, depending on the size of the compounds, it may be desirable to conjugate them to carrier proteins such as ovalbumin or serum albumin prior to their administration. Therapeutic formulations may be administered in any conventional dosage formulation. While it is possible for the active ingredient to be administered alone, it is preferable to present it as a pharmaceutical formulation. Formulations comprise at least one active ingredient, as defined above, together with one or more acceptable carriers thereof. Each carrier must be both pharmaceutically and

physiologically acceptable in the sense of being compatible with the other ingredients and not injurious to the patient. Formulations include those suitable for oral, rectal, nasal, or parenteral (including subcutaneous, intramuscular, intravenous and intradermal) administration. . The formulations may conveniently be presented in unit dosage form and may be prepared by any methods well known in the art of pharmacy. See, e.g., Gilman, et al. (eds. 1990) Goodman and Gilman's: The Pharmacological Bases of Therapeutics, 8th Ed., Pergamon Press; and Remington's Pharmaceutical Sciences (current edition), Mack Publishing Co., Easton, Penn.; Avis, et al. (eds. 1993) Pharmaceutical Dosage Forms: Parenteral Medications Dekker, NY; Lieberman, et al. (eds. 1990) Pharmaceutical Dosage Forms: Tablets Dekker, NY; and Lieberman, et al. (eds. 1990) Pharmaceutical Dosage Forms: Disperse Systems Dekker, NY. The therapy of this invention may be combined with or used in association with other therapeutic agents, particularly agonists or antagonists of other IL-1 family members.

IX. Ligands

The description of the Toll receptors herein provide means to identify ligands, as described above. Such ligand should bind specifically to the respective receptor with reasonably high affinity. Various constructs are made available which allow either labeling of the receptor to detect its ligand. For example, directly labeling DTLR, fusing onto it markers for secondary labeling, e.g., FLAG or other epitope tags, etc., will allow detection of receptor. This can be histological, as an affinity method for biochemical purification, or labeling or selection in an expression cloning approach. A two-hybrid selection system may also be applied making appropriate constructs with the available DTLR sequences. See, e.g., Fields and Song (1989) Nature 340:245-246.

Generally, descriptions of DTLRs will be analogously applicable to individual specific embodiments directed to DTLR2, DTLR3, DTLR4, DTLR5, DTLR6, DTLR7, DTLR8, DTLR9, and/or DTLR10 reagents and compositions.

The broad scope of this invention is best understood with reference to the following examples, which are not intended to limit the inventions to the specific embodiments.

EXAMPLES

I. General Methods

Some of the standard methods are described or referenced, e.g., in Maniatis, et al. (1982) Molecular Cloning, A Laboratory Manual, Cold Spring Harbor Laboratory, Cold Spring Harbor Press; Sambrook, et al. (1989) Molecular Cloning: A Laboratory Manual, (2d ed.), vols. 1-3, CSH Press, NY; Ausubel, et al., Biology, Greene Publishing Associates, Brooklyn, NY; or Ausubel, et al. (1987 and Supplements) Current Protocols in Molecular Biology, Greene/Wiley, New York. Methods for protein purification include such methods as ammonium sulfate precipitation, column chromatography, electrophoresis, centrifugation, crystallization, and others. See, e.g., Ausubel, et al. (1987 and periodic supplements); Coligan, et al. (ed. 1996) and periodic supplements, Current Protocols In Protein Science Greene/Wiley, New York; Deutscher (1990) "Guide to Protein Purification" in Methods in Enzymology, vol. 182, and other volumes in this series; and manufacturer's literature on use of protein purification products, e.g., Pharmacia, Piscataway, N.J., or Bio-Rad, Richmond, CA. Combination with recombinant techniques allow fusion to appropriate segments, e.g., to a FLAG sequence or an equivalent which can be fused via a protease-removable sequence. See, e.g., Hochuli (1989) Chemische Industrie 12:69-70; Hochuli (1990) "Purification of Recombinant Proteins with Metal Chelate Absorbent" in Setlow (ed.) Genetic Engineering, Principle and Methods 12:87-98, Plenum Press, N.Y.; and Crowe, et al. (1992) QIAexpress: The High Level Expression and Protein Purification System QUIAGEN, Inc., Chatsworth, CA.

Standard immunological techniques and assays are described, e.g., in Hertzenberg, et al. (eds. 1996) Weir's Handbook of Experimental Immunology vols. 1-4, Blackwell Science; Coligan (1991) Current Protocols in Immunology

Wiley/Greene, NY; and Methods in Enzymology volumes. 70, 73, 74, 84, 92, 93, 108, 116, 121, 132, 150, 162, and 163.

Assays for vascular biological activities are well known in the art. They will cover angiogenic and angiostatic activities in tumor, or other tissues, e.g., arterial smooth muscle proliferation (see, e.g., Koyoma, et al. (1996) Cell 87:1069-1078), monocyte adhesion to vascular epithelium (see McEvoy, et al. (1997) J. Exp. Med. 185:2069-2077), etc. See also Ross (1993) Nature 362:801-809; Rekhter and Gordon (1995) Am. J. Pathol. 147:668-677; Thyberg, et al. (1990) Atherosclerosis 10:966-990; and Gumbiner (1996) Cell 84:345-357.

Assays for neural cell biological activities are described, e.g., in Wouterlood (ed. 1995) Neuroscience Protocols modules 10, Elsevier; Methods in Neurosciences Academic Press; and Neuromethods Humana Press, Totowa, NJ. Methodology of developmental systems is described, e.g., in Meisami (ed.) Handbook of Human Growth and Developmental Biology CRC Press; and Chrispeels (ed.) Molecular Techniques and Approaches in Developmental Biology Interscience.

Computer sequence analysis is performed, e.g., using available software programs, including those from the GCG (U. Wisconsin) and GenBank sources. Public sequence databases were also used, e.g., from GenBank, NCBI, EMBO, and others. Determination of transmembrane and other important motifs may be predicted using such bioinformatics tools.

Many techniques applicable to IL-10 receptors may be applied to DTLRs, as described, e.g., in USSN 08/110,683 (IL-10 receptor), which is incorporated herein by reference for all purposes.

II. Novel Family of Human Receptors

Abbreviations: DTLR, DNAX Toll-like receptor; IL-1R, interleukin-1 receptor; TH, Toll homology; LRR, leucine-rich repeat; EST, expressed sequence tag; STS, sequence tagged site; FISH, fluorescence in situ hybridization.

The discovery of sequence homology between the cytoplasmic domains of Drosophila Toll and human interleukin-1 (IL-1) receptors has sown the conviction that both molecules trigger related signaling pathways tied to the nuclear translocation of Rel-type transcription factors. This conserved signaling scheme governs an evolutionarily ancient immune response in both insects and vertebrates. We report the molecular cloning of a novel class of putative human receptors with a protein architecture that is closely similar to Drosophila Toll in both intra- and extra-cellular segments. Five human Toll-like receptors, designated DTLRs 1-5, are likely the direct homologs of the fly molecule, and as such could constitute an important and unrecognized component of innate immunity in humans; intriguingly, the evolutionary retention of DTLRs in vertebrates may indicate another role, akin to Toll in the dorso-ventralization of the Drosophila embryo, as regulators of early morphogenetic patterning. Multiple tissue mRNA blots indicate markedly different patterns of expression for the human DTLRs. Using fluorescence in situ hybridization and Sequence-Tagged Site database analyses, we also show that the cognate DTLR genes reside on chromosomes 4 (DTLRs 1, 2, and 3), 9 (DTLR4), and 1 (DTLR5). Structure prediction of the aligned Toll-homology (TH) domains from varied insect and human DTLRs, vertebrate IL-1 receptors, and MyD88 factors, and plant disease resistance proteins, recognizes a parallel β/α fold with an acidic active site; a similar structure notably recurs in a class of response regulators broadly involved in transducing sensory information in bacteria.

The seeds of the morphogenetic gulf that so dramatically separates flies from humans are planted in familiar embryonic shapes and patterns, but give rise to very different cell complexities. DeRobertis and Sasai (1996) Nature 380:37-40; and Arendt and Nübler-Jung (1997) Mech. Develop. 61:7-21. This divergence of developmental plans between insects and vertebrates is choreographed by remarkably similar signaling pathways, underscoring a greater conservation of protein networks and biochemical mechanisms from unequal gene repertoires. Miklos and Rubin (1996) Cell 86:521-529; and Chothia (1994) Develop. 1994 Suppl., 27-33. A powerful way to chart the evolutionary design of these regulatory pathways is by inferring their likely molecular components (and biological functions) through interspecies comparisons of protein sequences and structures. Miklos and Rubin (1996) Cell 86:521-529; Chothia (1994) Develop. 1994 Suppl., 27-33 (3-5); and Banfi, et al. (1996) Nature Genet. 13:167-174.

A universally critical step in embryonic development is the specification of body axes, either born from innate asymmetries or triggered by external cues. DeRobertis and Sasai (1996) Nature 380:37-40; and Arendt and Nübler-Jung (1997) Mech. Develop. 61:7-21. As a model system, particular attention has been focused on the phylogenetic basis and cellular mechanisms of dorsoventral polarization. DeRobertis and Sasai (1996) Nature 380:37-40; and Arendt and Nübler-Jung (1997) Mech. Develop. 61:7-21. A prototype molecular strategy for this transformation has emerged from the Drosophila embryo, where the sequential action of a small number of genes results in a ventralizing gradient of the transcription factor Dorsal. St. Johnston and Nüsslein-Volhard (1992) Cell 68:201-219; and Morisato and Anderson (1995) Ann. Rev. Genet. 29:371-399.

This signaling pathway centers on Toll, a transmembrane receptor that transduces the binding of a maternally-secreted ventral factor, Spätzle, into the cytoplasmic engagement of Tube, an accessory molecule, and the activation of Pelle, a Ser/Thr kinase that catalyzes the dissociation of Dorsal from the inhibitor Cactus and allows migration of Dorsal to ventral nuclei (Morisato and Anderson (1995) Ann. Rev. Genet. 29:371-399; and Belvin and Anderson (1996) Ann. Rev. Cell Develop. Biol. 12:393-416. The Toll pathway also controls the induction of potent antimicrobial factors in the adult fly (Lemaitre, et al. (1996) Cell 86:973-983); this role in Drosophila immune defense strengthens mechanistic parallels to IL-1 pathways that govern a host of immune and inflammatory responses in vertebrates. Belvin and Anderson (1996) Ann. Rev. Cell Develop. Biol. 12:393-416; and Wasserman (1993) Molec. Biol. Cell 4:767-771. A Toll-related cytoplasmic domain in IL-1 receptors directs the binding of a Pelle-like kinase, IRAK, and the activation of a latent NF-κB/I-κB complex that mirrors the embrace of Dorsal and Cactus. Belvin and Anderson (1996) Ann. Rev. Cell Develop. Biol. 12:393-416; and Wasserman (1993) Molec. Biol. Cell 4:767-771.

We describe the cloning and molecular characterization of four new Toll-like molecules in humans, designated DTLRs 2-5 (following Chiang and Beachy (1994) Mech. Develop. 47:225-239), that reveal a receptor family more closely tied to Drosophila Toll homologs than to vertebrate IL-1 receptors. The DTLR sequences are derived from human ESTs; these partial cDNAs were used to draw complete expression profiles in human tissues for the five DTLRs, map the chromosomal locations of cognate genes, and narrow the choice of cDNA libraries for full-length cDNA retrievals. Spurred by other efforts (Banfi, et al. (1996) Nature Genet. 13:167-174; and Wang, et al. (1996) J. Biol. Chem. 271:4468-4476), we are assembling,

by structural conservation and molecular parsimony, a biological system in humans that is the counterpart of a compelling regulatory scheme in Drosophila. In addition, a biochemical mechanism driving Toll signaling is suggested by the proposed tertiary fold of the Toll-homology (TH) domain, a core module shared by DTLRs, a broad family of IL-1 receptors, mammalian MyD88 factors and plant disease resistance proteins. Mitcham, et al. (1996) J. Biol. Chem. 271:5777-5783; and Hardiman, et al. (1996) Oncogene 13:2467-2475. We propose that a signaling route coupling morphogenesis and primitive immunity in insects, plants, and animals (Belvin and Anderson (1996) Ann. Rev. Cell Develop. Biol. 12:393-416; and Wilson, et al. (1997) Curr. Biol. 7:175-178) may have roots in bacterial two-component pathways.

Computational Analysis.

Human sequences related to insect DTLRs were identified from the EST database (dbEST) at the National Center for Biotechnology Information (NCBI) using the BLAST server (Altschul, et al. (1994) Nature Genet. 6:119-129). More sensitive pattern- and profile-based methods (Bork and Gibson (1996) Meth. Enzymol. 266:162-184) were used to isolate the signaling domains of the DTLR family that are shared with vertebrate and plant proteins present in nonredundant databases. The progressive alignment of DTLR intra- or extracellular domain sequences was carried out by ClustalW (Thompson, et al. (1994) Nucleic Acids Res. 22:4673-4680); this program also calculated the branching order of aligned sequences by the Neighbor-Joining algorithm (5000 bootstrap replications provided confidence values for the tree groupings).

Conserved alignment patterns, discerned at several degrees of stringency, were drawn by the Consensus program (internet URL http://www.bork.embl-heidelberg.de/Alignment/ consensus.html). The PRINTS

library of protein fingerprints (http://www.biochem.ucl.ac.uk/bsm/dbbrowser/PRINTS/PRINTS.html) (Attwood, et al. (1997) Nucleic Acids Res. 25:212-217) reliably identified the myriad leucine-rich repeats (LRRs) present in the extracellular segments of DTLRs with a compound motif (PRINTS code Leurichrpt) that flexibly matches N- and C-terminal features of divergent LRRs. Two prediction algorithms whose three-state accuracy is above 72% were used to derive a consensus secondary structure for the intracellular domain alignment, as a bridge to fold recognition efforts (Fischer, et al. (1996) FASEB J. 10:126-136). Both the neural network program PHD (Rost and Sander (1994) Proteins 19:55-72) and the statistical prediction method DSC (King and Sternberg (1996) Protein Sci. 5:2298-2310) have internet servers (URLs http://www.embl-heidelberg.de/predictprotein/phd_pred.html and http://bonsai.lif.icnet.uk/bmm/dsc/dsc_read_align.html, respectively). The intracellular region encodes the THD region discussed, e.g., in Hardiman, et al. (1996) Oncogene 13:2467-2475; and Rock, et al. (1998) Proc. Nat'l Acad. Sci. USA 95:588-593, each of which is incorporated herein by reference. This domain is very important in the mechanism of signaling by the receptors, which transfers a phosphate group to a substrate.

Cloning of full-length human DTLR cDNAs.

PCR primers derived from the Toll-like Humrsc786 sequence (GenBank accession code D13637) (Nomura, et al. (1994) DNA Res. 1:27-35) were used to probe a human erythroleukemic, TF-1 cell line-derived cDNA library (Kitamura, et al. (1989) Blood 73:375-380) to yield the DTLR1 cDNA sequence. The remaining DTLR sequences were flagged from dbEST, and the relevant EST clones obtained from the I.M.A.G.E. consortium (Lennon, et al. (1996) Genomics 33:151-152) via Research Genetics (Huntsville,

AL): CloneID#'s 80633 and 117262 (DTLR2), 144675 (DTLR3), 202057 (DTLR4) and 277229 (DTLR5). Full length cDNAs for human DTLRs 2-4 were cloned by DNA hybridization screening of λgt10 phage, human adult lung, placenta, and fetal liver 5'-Stretch Plus cDNA libraries (Clontech), respectively; the DTLR5 sequence is derived from a human multiple-sclerosis plaque EST. All positive clones were sequenced and aligned to identify individual DTLR ORFs: DTLR1 (2366 bp clone, 786 aa ORF), DTLR2 (2600 bp, 784 aa), DTLR3 (3029 bp, 904 aa), DTLR4 (3811 bp, 879 aa) and DTLR5 (1275 bp, 370 aa). Similar methods are used for DTLRs 6-10. Probes for DTLR3 and DTLR4 hybridizations were generated by PCR using human placenta (Stratagene) and adult liver (Clontech) cDNA libraries as templates, respectively; primer pairs were derived from the respective EST sequences. PCR reactions were conducted using T. aquaticus Taqplus DNA polymerase (Stratagene) under the following conditions: 1 x (94° C, 2 min) 30 x (55° C, 20 sec; 72° C 30 sec; 94° C 20 sec), 1 x (72° C, 8 min). For DTLR2 full-length cDNA screening, a 900 bp fragment generated by EcoRI/XbaI digestion of the first EST clone (ID# 80633) was used as a probe.

mRNA blots and chromosomal localization.

Human multiple tissue (Cat# 1, 2) and cancer cell line blots (Cat# 7757-1), containing approximately 2 μg of poly(A)$^+$ RNA per lane, were purchased from Clontech (Palo Alto, CA). For DTLRs 1-4, the isolated full-length cDNAs served as probes, for DTLR5 the EST clone (ID #277229) plasmid insert was used. Briefly, the probes were radiolabeled with [α-^{32}P] dATP using the Amersham Rediprime random primer labeling kit (RPN1633). Prehybridization and hybridizations were performed at 65° C in 0.5 M Na_2HPO_4, 7% SDS, 0.5 M EDTA (pH 8.0). All stringency washes were conducted at 65° C with two initial washes in 2 x SSC, 0.1% SDS for 40 min followed by a

subsequent wash in 0.1 x SSC, 0.1% SDS for 20 min. Membranes were then exposed at -70° C to X-Ray film (Kodak) in the presence of intensifying screens. More detailed studies by cDNA library Southerns (14) were performed with selected human DTLR clones to examine their expression in hemopoietic cell subsets.

Human chromosomal mapping was conducted by the method of fluorescence in situ hybridization (FISH) as described in Heng and Tsui (1994) Meth. Molec. Biol. 33:109-122, using the various full-length (DTLRs 2-4) or partial (DTLR5) cDNA clones as probes. These analyses were performed as a service by SeeDNA Biotech Inc. (Ontario, Canada). A search for human syndromes (or mouse defects in syntenic loci) associated with the mapped DTLR genes was conducted in the Dysmorphic Human-Mouse Homology Database by internet server (http://www.hgmp.mrc.ac.uk/DHMHD/ hum_chrome1.html). Similar methods nare applicable to DTLRs 6-10.

Conserved architecture of insect and human DTLR ectodomains.

The Toll family in Drosophila comprises at least four distinct gene products: Toll, the prototype receptor involved in dorsoventral patterning of the fly embryo (Morisato and Anderson (1995) Ann. Rev. Genet. 29:371-399) and a second named '18 Wheeler' (18w) that may also be involved in early embryonic development (Chiang and Beachy (1994) Mech. Develop. 47:225-239; Eldon, et al. (1994) Develop. 120:885-899); two additional receptors are predicted by incomplete, Toll-like ORFs downstream of the male-specific-transcript (Mst) locus (GenBank code X67703) or encoded by the 'sequence-tagged-site' (STS) Dm2245 (GenBank code G01378) (Mitcham, et al. (1996) J. Biol. Chem. 271:5777-5783). The extracellular segments of Toll and 18w are distinctively composed of imperfect, ~24 amino acid LRR motifs (Chiang and Beachy (1994) Mech. Develop.

47:225-239; and Eldon, et al. (1994) Develop. 120:885-899). Similar tandem arrays of LRRs commonly form the adhesive antennae of varied cell surface molecules and their generic tertiary structure is presumed to mimic the horseshoe-shaped cradle of a ribonuclease inhibitor fold, where seventeen LRRs show a repeating β/α-hairpin, 28 residue motif (Buchanan and Gay (1996) Prog. Biophys. Molec. Biol. 65:1-44). The specific recognition of Spätzle by Toll may follow a model proposed for the binding of cystine-knot fold glycoprotein hormones by the multi-LRR ectodomains of serpentine receptors, using the concave side of the curved β-sheet (Kajava, et al. (1995) Structure 3:867-877); intriguingly, the pattern of cysteines in Spätzle, and an orphan Drosophila ligand, Trunk, predict a similar cystine-knot tertiary structure (Belvin and Anderson (1996) Ann. Rev. Cell Develop. Biol. 12:393-416; and Casanova, et al. (1995) Genes Develop. 9:2539-2544).

The 22 and 31 LRR ectodomains of Toll and 18w, respectively (the Mst ORF fragment displays 16 LRRs), are most closely related to the comparable 18, 19, 24, and 22 LRR arrays of DTLRs 1-4 (the incomplete DTLR5 chain presently includes four membrane-proximal LRRs) by sequence and pattern analysis (Altschul, et al. (1994) Nature Genet. 6:119-129; and Bork and Gibson (1996) Meth. Enzymol. 266:162-184) (Fig. 1). However, a striking difference in the human DTLR chains is the common loss of a ~90 residue cysteine-rich region that is variably embedded in the ectodomains of Toll, 18w and the Mst ORF (distanced four, six and two LRRs, respectively, from the membrane boundary). These cysteine clusters are bipartite, with distinct 'top' (ending an LRR) and 'bottom' (stacked atop an LRR) halves (Chiang and Beachy (1994) Mech. Develop. 47:225-239; Eldon, et al. (1994) Develop. 120:885-899; and Buchanan and Gay (1996) Prog. Biophys. Molec. Biol. 65:1-44); the 'top' module recurs in

both Drosophila and human DTLRs as a conserved juxtamembrane spacer (Fig. 1). We suggest that the flexibly located cysteine clusters in Drosophila receptors (and other LRR proteins), when mated 'top' to 'bottom', form a compact module with paired termini that can be inserted between any pair of LRRs without altering the overall fold of DTLR ectodomains; analogous 'extruded' domains decorate the structures of other proteins (Russell (1994) Protein Engin. 7:1407-1410).

Molecular design of the TH signaling domain.

Sequence comparison of Toll and IL-1 type-I (IL-1R1) receptors has disclosed a distant resemblance of a ~200 amino acid cytoplasmic domain that presumably mediates signaling by similar Rel-type transcription factors. Belvin and Anderson (1996) Ann. Rev. Cell Develop. Biol. 12:393-416; and (Belvin and Anderson (1996) Ann. Rev. Cell Develop. Biol. 12:393-416; and Wasserman (1993) Molec. Biol. Cell 4:767-771). More recent additions to this functional paradigm include a pair of plant disease resistance proteins from tobacco and flax that feature an N-terminal TH module followed by nucleotide-binding (NTPase) and LRR segments (Wilson, et al. (1997) Curr. Biol. 7:175-178); by contrast, a 'death domain' precedes the TH chain of MyD88, an intracellular myeloid differentiation marker (Mitcham, et al. (1996) J. Biol. Chem. 271:5777-5783; and Hardiman, et al. (1996) Oncogene 13:2467-2475) (Fig. 1). New IL-1-type receptors include IL-1R3, an accessory signaling molecule, and orphan receptors IL-1R4 (also called ST2/Fit-1/T1), IL-1R5 (IL-1R-related protein), and IL-1R6 (IL-1R-related protein-2) (Mitcham, et al. (1996) J. Biol. Chem. 271:5777-5783;Hardiman, et al. (1996) Oncogene 13:2467-2475). With the new human DTLR sequences, we have sought a structural definition of this evolutionary thread by analyzing the conformation of the common TH module: ten blocks of

conserved sequence comprising 128 amino acids form the minimal TH domain fold; gaps in the alignment mark the likely location of sequence and length-variable loops (Fig. 2A-2B).

Two prediction algorithms that take advantage of the patterns of conservation and variation in multiply aligned sequences, PHD (Rost and Sander (1994) Proteins 19:55-72) and DSC (King and Sternberg (1996) Protein Sci. 5:2298-2310), produced strong, concordant results for the TH signaling module (Fig. 2A-2B). Each block contains a discrete secondary structural element: the imprint of alternating β-strands (labeled A-E) and α-helices (numbered 1-5) is diagnostic of a β/α-class fold with α-helices on both faces of a parallel β-sheet. Hydrophobic β-strands A, C and D are predicted to form 'interior' staves in the β-sheet, while the shorter, amphipathic β-strands B and E resemble typical 'edge' units (Fig. 2A-2B). This assignment is consistent with a strand order of B-A-C-D-E in the core β-sheet (Fig. 2C); fold comparison ('mapping') and recognition ('threading') programs (Fischer, et al. (1996) FASEB J. 10:126-136) strongly return this doubly wound β/α topology. A surprising, functional prediction of this outline structure for the TH domain is that many of the conserved, charged residues in the multiple alignment map to the C-terminal end of the β-sheet: residue Asp16 (block numbering scheme - Fig. 2A-2B) at the end of βA, Arg39 and Asp40 following βB, Glu75 in the first turn of α3, and the more loosely conserved Glu/Asp residues in the βD-α4 loop, or after βE (Fig. 2A-2B). The location of four other conserved residues (Asp7, Glu28, and the Arg57-Arg/Lys58 pair) is compatible with a salt bridge network at the opposite, N-terminal end of the β-sheet (Fig. 2A-2B). Alignment of the other DTLR embodiments exhibit similar features, and peptide segments comprising these feataures, e.g., 20 amino acid segments containing them, are particularly important.

Signaling function depends on the structural integrity of the TH domain. Inactivating mutations or deletions within the module boundaries (Fig. 2A-2B) have been catalogued for IL-1R1 and Toll. Heguy, et al. (1992) J. Biol. Chem. 267:2605-2609; Croston, et al. (1995) J. Biol. Chem. 270;16514-16517; Schneider, et al. (1991) Genes Develop. 5:797-807; Norris and Manley. (1992) Genes Develop. 6:1654-1667; Norris and Manley (1995) Genes Develop. 9:358-369; and Norris and Manley (1996) Genes Develop. 10:862-872. The human DTLR1-5 chains extending past the minimal TH domain (8, 0, 6, 22 and 18 residue lengths, respectively) are most closely similar to the stubby, 4 aa 'tail' of the Mst ORF. Toll and 18w display unrelated 102 and 207 residue tails (Fig. 2A-2B) that may negatively regulate the signaling of the fused TH domains. Norris and Manley (1995) Genes Develop. 9:358-369; and Norris and Manley (1996) Genes Develop. 10:862-872.

The evolutionary relationship between the disparate proteins that carry the TH domain can best be discerned by a phylogenetic tree derived from the multiple alignment (Fig. 3). Four principal branches segregate the plant proteins, the MyD88 factors, IL-1 receptors, and Toll-like molecules; the latter branch clusters the Drosophila and human DTLRs.

Chromosomal dispersal of human DTLR genes.

In order to investigate the genetic linkage of the nascent human DTLR gene family, we mapped the chromosomal loci of four of the five genes by FISH (Fig. 4). The DTLR1 gene has previously been charted by the human genome project: an STS database locus (dbSTS accession number G06709, corresponding to STS WI-7804 or SHGC-12827) exists for the Humrsc786 cDNA (Nomura, et al. (1994) DNA Res. 1:27-35) and fixes the gene to chromosome 4 marker interval D4S1587-D42405 (50-56 cM) circa 4p14. This assignment has recently been corroborated by FISH

analysis. Taguchi, et al. (1996) Genomics 32:486-488. In the present work, we reliably assign the remaining DTLR genes to loci on chromosome 4q32 (DTLR2), 4q35 (DTLR3), 9q32-33 (DTLR4) and 1q33.3 (DTLR5). During the course of this work, an STS for the parent DTLR2 EST (cloneID # 80633) has been generated (dbSTS accession number T57791 for STS SHGC-33147) and maps to the chromosome 4 marker interval D4S424-D4S1548 (143-153 cM) at 4q32 -in accord with our findings. There is a ~50 cM gap between DTLR2 and DTLR3 genes on the long arm of chromosome 4.

DTLR genes are differentially expressed.

Both Toll and 18w have complex spatial and temporal patterns of expression in Drosophila that may point to functions beyond embryonic patterning. St. Johnston and Nüsslein-Volhard (1992) Cell 68:201-219; Morisato and Anderson (1995) Ann. Rev. Genet. 29:371-399; Belvin and Anderson (1996) Ann. Rev. Cell Develop. Biol. 12:393-416; Lemaitre, et al. (1996) Cell 86:973-983; Chiang and Beachy (1994) Mech. Develop. 47:225-239; and Eldon, et al. (1994) Develop. 120:885-899. We have examined the spatial distribution of DTLR transcripts by mRNA blot analysis with varied human tissue and cancer cell lines using radiolabeled DTLR cDNAs (Fig. 5). DTLR1 is found to be ubiquitously expressed, and at higher levels than the other receptors. Presumably reflecting alternative splicing, 'short' 3.0 kB and 'long' 8.0 kB DTLR1 transcript forms are present in ovary and spleen, respectively (Fig. 5, panels A and B). A cancer cell mRNA panel also shows the prominent overexpression of DTLR1 in a Burkitt's Lymphoma Raji cell line (Fig. 5, panel C). DTLR2 mRNA is less widely expressed than DTLR1, with a 4.0 kB species detected in lung and a 4.4 kB transcript evident in heart, brain and muscle. The tissue distribution pattern of DTLR3 echoes that of DTLR2 (Fig. 5, panel E). DTLR3 is also present as two major

transcripts of approximately 4.0 and 6.0 kB in size, and the highest levels of expression are observed in placenta and pancreas. By contrast, DTLR4 and DTLR5 messages appear to be extremely tissue-specific. DTLR4 was detected only in placenta as a single transcript of ~7.0 kB in size. A faint 4.0 kB signal was observed for DTLR5 in ovary and peripheral blood monocytes.

Components of an evolutionarily ancient regulatory system.

The original molecular blueprints and divergent fates of signaling pathways can be reconstructed by comparative genomic approaches. Miklos and Rubin (1996) Cell 86:521-529; Chothia (1994) Develop. 1994 Suppl., 27-33; Banfi, et al. (1996) Nature Genet. 13:167-174; and Wang, et al. (1996) J. Biol. Chem. 271:4468-4476. We have used this logic to identify an emergent gene family in humans, encoding five receptor paralogs at present, DTLRs 1-5, that are the direct evolutionary counterparts of a Drosophila gene family headed by Toll (Figs. 1-3). The conserved architecture of human and fly DTLRs, conserved LRR ectodomains and intracellular TH modules (Fig. 1), intimates that the robust pathway coupled to Toll in Drosophila (6, 7) survives in vertebrates. The best evidence borrows from a reiterated pathway: the manifold IL-1 system and its repertoire of receptor-fused TH domains, IRAK, NF-κB and I-κB homologs (Belvin and Anderson (1996) Ann. Rev. Cell Develop. Biol. 12:393-416; Wasserman (1993) Molec. Biol. Cell 4:767-771; Hardiman, et al. (1996) Oncogene 13:2467-2475; and Cao, et al. (1996) Science 271:1128-1131); a Tube-like factor has also been characterized. It is not known whether DTLRs can productively couple to the IL-1R signaling machinery, or instead, a parallel set of proteins is used. Differently from IL-1 receptors, the LRR cradle of human DTLRs is predicted to retain an affinity for Spätzle/Trunk-related

cystine-knot factors; candidate DTLR ligands (called PENs) that fit this mold have been isolated.

Biochemical mechanisms of signal transduction can be gauged by the conservation of interacting protein folds in a pathway. Miklos and Rubin (1996) Cell 86:521-529; Chothia (1994) Develop. 1994 Suppl., 27-33. At present, the Toll signaling paradigm involves some molecules whose roles are narrowly defined by their structures, actions or fates: Pelle is a Ser/Thr kinase (phosphorylation), Dorsal is an NF-κB-like transcription factor (DNA-binding) and Cactus is an ankyrin-repeat inhibitor (Dorsal binding, degradation). Belvin and Anderson (1996) Ann. Rev. Cell Develop. Biol. 12:393-416. By contrast, the functions of the Toll TH domain and Tube remain enigmatic. Like other cytokine receptors (Heldin (1995) Cell 80:213-223), ligand-mediated dimerization of Toll appears to be the triggering event: free cysteines in the juxtamembrane region of Toll create constitutively active receptor pairs (Schneider, et al. (1991) Genes Develop. 5:797-807), and chimeric Torso-Toll receptors signal as dimers (Galindo, et al. (1995) Develop. 121:2209-2218); yet, severe truncations or wholesale loss of the Toll ectodomain results in promiscuous intracellular signaling (Norris and Manley (1995) Genes Develop. 9:358-369; and Winans and Hashimoto (1995) Molec. Biol. Cell 6:587-596), reminiscent of oncogenic receptors with catalytic domains (Heldin (1995) Cell 80:213-223). Tube is membrane-localized, engages the N-terminal (death) domain of Pelle and is phosphorylated, but neither Toll-Tube or Toll-Pelle interactions are registered by two-hybrid analysis (Galindo, et al. (1995) Develop. 121:2209-2218; and Groβhans, et al. (1994) Nature 372:563-566); this latter result suggests that the conformational 'state' of the Toll TH domain somehow affects factor recruitment. Norris and Manley (1996) Genes Develop. 10:862-872; and Galindo, et al. (1995) Develop. 121:2209-2218.

At the heart of these vexing issues is the structural nature of the Toll TH module. To address this question, we have taken advantage of the evolutionary diversity of TH sequences from insects, plants and vertebrates, incorporating the human DTLR chains, and extracted the minimal, conserved protein core for structure prediction and fold recognition (Fig. 2). The strongly predicted $(\beta/\alpha)_5$ TH domain fold with its asymmetric cluster of acidic residues is topologically identical to the structures of response regulators in bacterial two-component signaling pathways (Volz (1993) Biochemistry 32:11741-11753; and Parkinson (1993) Cell 73:857-871) (Fig. 2A-2C). The prototype chemotaxis regulator CheY transiently binds a divalent cation in an 'aspartate pocket' at the C-end of the core β-sheet; this cation provides electrostatic stability and facilitates the activating phosphorylation of an invariant Asp. Volz (1993) Biochemistry 32:11741-11753. Likewise, the TH domain may capture cations in its acidic nest, but activation, and downstream signaling, could depend on the specific binding of a negatively charged moiety: anionic ligands can overcome intensely negative binding-site potentials by locking into precise hydrogen-bond networks. Ledvina, et al. (1996) Proc. Natl. Acad. Sci. USA 93:6786-6791. Intriguingly, the TH domain may not simply act as a passive scaffold for the assembly of a Tube/Pelle complex for Toll, or homologous systems in plants and vertebrates, but instead actively participate as a true conformational trigger in the signal transducing machinery. Perhaps explaining the conditional binding of a Tube/Pelle complex, Toll dimerization could promote unmasking, by regulatory receptor tails (Norris and Manley (1995) Genes Develop. 9:358-369; Norris and Manley (1996) Genes Develop. 10:862-872), or binding by small molecule activators of the TH pocket. However, 'free' TH modules inside the cell (Norris and Manley (1995) Genes Develop. 9:358-369; Winans and Hashimoto

(1995) Molec. Biol. Cell 6:587-596) could act as catalytic, CheY-like triggers by activating and docking with errant Tube/Pelle complexes.

Morphogenetic receptors and immune defense.

The evolutionary link between insect and vertebrate immune systems is stamped in DNA: genes encoding antimicrobial factors in insects display upstream motifs similar to acute phase response elements known to bind NF-κB transcription factors in mammals. Hultmark (1993) Trends Genet. 9:178-183. Dorsal, and two Dorsal-related factors, Dif and Relish, help induce these defense proteins after bacterial challenge (Reichhart, et al. (1993) C. R. Acad. Sci. Paris 316:1218-1224; Ip, et al. (1993) Cell 75:753-763; and Dushay, et al. (1996) Proc. Natl. Acad. Sci. USA 93:10343-10347); Toll, or other DTLRs, likely modulate these rapid immune responses in adult Drosophila (Lemaitre, et al. (1996) Cell 86:973-983; and Rosetto, et al. (1995) Biochem. Biophys. Res. Commun. 209:111-116). These mechanistic parallels to the IL-1 inflammatory response in vertebrates are evidence of the functional versatility of the Toll signaling pathway, and suggest an ancient synergy between embryonic patterning and innate immunity (Belvin and Anderson (1996) Ann. Rev. Cell Develop. Biol. 12:393-416; Lemaitre, et al. (1996) Cell 86:973-983; Wasserman (1993) Molec. Biol. Cell 4:767-771; Wilson, et al. (1997) Curr. Biol. 7:175-178; Hultmark (1993) Trends Genet. 9:178-183; Reichhart, et al. (1993) C. R. Acad. Sci. Paris 316:1218-1224; Ip, et al. (1993) Cell 75:753-763; Dushay, et al. (1996) Proc. Natl. Acad. Sci. USA 93:10343-10347; Rosetto, et al. (1995) Biochem. Biophys. Res. Commun. 209:111-116; Medzhitov and Janeway (1997) Curr. Opin. Immunol. 9:4-9; and Medzhitov and Janeway (1997) Curr. Opin. Immunol. 9:4-9). The closer homology of insect and human DTLR proteins invites an even stronger overlap of biological functions that supersedes

the purely immune parallels to IL-1 systems, and lends potential molecular regulators to dorso-ventral and other transformations of vertebrate embryos. DeRobertis and Sasai (1996) Nature 380:37-40; and Arendt and Nübler-Jung (1997) Mech. Develop. 61:7-21.

The present description of an emergent, robust receptor family in humans mirrors the recent discovery of the vertebrate Frizzled receptors for Wnt patterning factors. Wang, et al. (1996) J. Biol. Chem. 271:4468-4476. As numerous other cytokine-receptor systems have roles in early development (Lemaire and Kodjabachian (1996) Trends Genet. 12:525-531), perhaps the distinct cellular contexts of compact embryos and gangly adults simply result in familiar signaling pathways and their diffusible triggers having different biological outcomes at different times, e.g., morphogenesis versus immune defense for DTLRs. For insect, plant, and human Toll-related systems (Hardiman, et al. (1996) Oncogene 13:2467-2475; Wilson, et al. (1997) Curr. Biol. 7:175-178), these signals course through a regulatory TH domain that intriguingly resembles a bacterial transducing engine (Parkinson (1993) Cell 73:857-871).

In particular, the DTLR6 exhibits structural features which establish its membership in the family. Moreover, members of the family have been implicated in a number of significant developmental disease conditions and with function of the innate immune system. In particular, the DTLR6 has been mapped to the X chromosome to a location which is a hot spot for major developmental abnormalities. See, e.g., The Sanger Center: human X chromosome website http://www.sanger.ac.uk/HGP/ChrX/index.shtml; and the Baylor College of Medicine Human Genome Sequencing website http://gc.bcm.tmc.edu:8088/cgi-bin/seq/home.

The accession number for the deposited PAC is AC003046. This accession number contains sequence from two PACs: RPC-164K3 and RPC-263P4. These two PAC

sequences mapped on human chromosome Xp22 at the Baylor web site between STS markers DXS704 and DXS7166. This region is a "hot spot" for severe developmental abnormalities.

III. Amplification of DTLR fragment by PCR

Two appropriate primer sequences are selected (see Tables 1 through 10). RT-PCR is used on an appropriate mRNA sample selected for the presence of message to produce a partial or full length cDNA, e.g., a sample which expresses the gene. See, e.g., Innis, et al. (eds. 1990) PCR Protocols: A Guide to Methods and Applications Academic Press, San Diego, CA; and Dieffenbach and Dveksler (eds. 1995) PCR Primer: A Laboratory Manual Cold Spring Harbor Press, CSH, NY. Such will allow determination of a useful sequence to probe for a full length gene in a cDNA library. The DTLR6 is a contiguous sequence in the genome, which may suggest that the other DTLRs are also. Thus, PCR on genomic DNA may yield full length contiguous sequence, and chromosome walking methodology would then be applicable. Alternatively, sequence databases will contain sequence corresponding to portions of the described embodiments, or closely related forms, e.g., alternative splicing, etc. Expression cloning techniques also may be applied on cDNA libraries.

IV. Tissue distribution of DTLRs

Message for each gene encoding these DTLRs has been detected. See Figures 5A-5F. Other cells and tissues will be assayed by appropriate technology, e.g., PCR, immunoassay, hybridization, or otherwise. Tissue and organ cDNA preparations are available, e.g., from Clontech, Mountain View, CA. Identification of sources of natural expression are useful, as described.

Southern Analysis: DNA (5 µg) from a primary amplifie cDNA library is digested with appropriate restriction

enzymes to release the inserts, run on a 1% agarose gel and transferred to a nylon membrane (Schleicher and Schuell, Keene, NH).

Samples for human mRNA isolation would typically include, e.g.: peripheral blood mononuclear cells (monocytes, T cells, NK cells, granulocytes, B cells), resting (T100); peripheral blood mononuclear cells, activated with anti-CD3 for 2, 6, 12 h pooled (T101); T cell, TH0 clone Mot 72, resting (T102); T cell, TH0 clone Mot 72, activated with anti-CD28 and anti-CD3 for 3, 6, 12 h pooled (T103); T cell, TH0 clone Mot 72, anergic treated with specific peptide for 2, 7, 12 h pooled (T104); T cell, TH1 clone HY06, resting (T107); T cell, TH1 clone HY06, activated with anti-CD28 and anti-CD3 for 3, 6, 12 h pooled (T108); T cell, TH1 clone HY06, anergic treated with specific peptide for 2, 6, 12 h pooled (T109); T cell, TH2 clone HY935, resting (T110); T cell, TH2 clone HY935, activated with anti-CD28 and anti-CD3 for 2, 7, 12 h pooled (T111); T cells CD4+CD45RO- T cells polarized 27 days in anti-CD28, IL-4, and anti IFN-γ, TH2 polarized, activated with anti-CD3 and anti-CD28 4 h (T116); T cell tumor lines Jurkat and Hut78, resting (T117); T cell clones, pooled AD130.2, Tc783.12, Tc783.13, Tc783.58, Tc782.69, resting (T118); T cell random γδ T cell clones, resting (T119); Splenocytes, resting (B100); Splenocytes, activated with anti-CD40 and IL-4 (B101); B cell EBV lines pooled WT49, RSB, JY, CVIR, 721.221, RM3, HSY, resting (B102); B cell line JY, activated with PMA and ionomycin for 1, 6 h pooled (B103); NK 20 clones pooled, resting (K100); NK 20 clones pooled, activated with PMA and ionomycin for 6 h (K101); NKL clone, derived from peripheral blood of LGL leukemia patient, IL-2 treated (K106); NK cytotoxic clone 640-A30-1, resting (K107); hematopoietic precursor line TF1, activated with PMA and ionomycin for 1, 6 h pooled (C100); U937 premonocytic line, resting (M100); U937 premonocytic line, activated

with PMA and ionomycin for 1, 6 h pooled (M101); elutriated monocytes, activated with LPS, IFNγ, anti-IL-10 for 1, 2, 6, 12, 24 h pooled (M102); elutriated monocytes, activated with LPS, IFNγ, IL-10 for 1, 2, 6, 12, 24 h pooled (M103); elutriated monocytes, activated with LPS, IFNγ, anti-IL-10 for 4, 16 h pooled (M106); elutriated monocytes, activated with LPS, IFNγ, IL-10 for 4, 16 h pooled (M107); elutriated monocytes, activated LPS for 1 h (M108); elutriated monocytes, activated LPS for 6 h (M109); DC 70% CD1a+, from CD34+ GM-CSF, TNFα 12 days, resting (D101); DC 70% CD1a+, from CD34+ GM-CSF, TNFα 12 days, activated with PMA and ionomycin for 1 hr (D102); DC 70% CD1a+, from CD34+ GM-CSF, TNFα 12 days, activated with PMA and ionomycin for 6 hr (D103); DC 95% CD1a+, from CD34+ GM-CSF, TNFα 12 days FACS sorted, activated with PMA and ionomycin for 1, 6 h pooled (D104); DC 95% CD14+, ex CD34+ GM-CSF, TNFα 12 days FACS sorted, activated with PMA and ionomycin 1, 6 hr pooled (D105); DC CD1a+ CD86+, from CD34+ GM-CSF, TNFα 12 days FACS sorted, activated with PMA and ionomycin for 1, 6 h pooled (D106); DC from monocytes GM-CSF, IL-4 5 days, resting (D107); DC from monocytes GM-CSF, IL-4 5 days, resting (D108); DC from monocytes GM-CSF, IL-4 5 days, activated LPS 4, 16 h pooled (D109); DC from monocytes GM-CSF, IL-4 5 days, activated TNFα, monocyte supe for 4, 16 h pooled (D110); leiomyoma L11 benign tumor (X101); normal myometrium M5 (O115); malignant leiomyosarcoma GS1 (X103); lung fibroblast sarcoma line MRC5, activated with PMA and ionomycin for 1, 6 h pooled (C101); kidney epithelial carcinoma cell line CHA, activated with PMA and ionomycin for 1, 6 h pooled (C102); kidney fetal 28 wk male (O100); lung fetal 28 wk male (O101); liver fetal 28 wk male (O102); heart fetal 28 wk male (O103); brain fetal 28 wk male (O104); gallbladder fetal 28 wk male (O106); small intestine fetal 28 wk male (O107); adipose tissue fetal 28 wk male (O108); ovary fetal 25 wk female (O109); uterus fetal 25 wk female

(O110); testes fetal 28 wk male (O111); spleen fetal 28 wk male (O112); adult placenta 28 wk (O113); and tonsil inflamed, from 12 year old (X100).

Samples for mouse mRNA isolation can include, e.g.: resting mouse fibroblastic L cell line (C200); Braf:ER (Braf fusion to estrogen receptor) transfected cells, control (C201); T cells, TH1 polarized (Mel14 bright, CD4+ cells from spleen, polarized for 7 days with IFN-γ and anti IL-4; T200); T cells, TH2 polarized (Mel14 bright, CD4+ cells from spleen, polarized for 7 days with IL-4 and anti-IFN-γ; T201); T cells, highly TH1 polarized (see Openshaw, et al. (1995) J. Exp. Med. 182:1357-1367; activated with anti-CD3 for 2, 6, 16 h pooled; T202); T cells, highly TH2 polarized (see Openshaw, et al. (1995) J. Exp. Med. 182:1357-1367; activated with anti-CD3 for 2, 6, 16 h pooled; T203); CD44- CD25+ pre T cells, sorted from thymus (T204); TH1 T cell clone D1.1, resting for 3 weeks after last stimulation with antigen (T205); TH1 T cell clone D1.1, 10 µg/ml ConA stimulated 15 h (T206); TH2 T cell clone CDC35, resting for 3 weeks after last stimulation with antigen (T207); TH2 T cell clone CDC35, 10 µg/ml ConA stimulated 15 h (T208); Mel14+ naive T cells from spleen, resting (T209); Mel14+ T cells, polarized to Th1 with IFN-γ/IL-12/anti-IL-4 for 6, 12, 24 h pooled (T210); Mel14+ T cells, polarized to Th2 with IL-4/anti-IFN-γ for 6, 13, 24 h pooled (T211); unstimulated mature B cell leukemia cell line A20 (B200); unstimulated B cell line CH12 (B201); unstimulated large B cells from spleen (B202); B cells from total spleen, LPS activated (B203); metrizamide enriched dendritic cells from spleen, resting (D200); dendritic cells from bone marrow, resting (D201); monocyte cell line RAW 264.7 activated with LPS 4 h (M200); bone-marrow macrophages derived with GM and M-CSF (M201); macrophage cell line J774, resting (M202); macrophage cell line J774 + LPS + anti-IL-10 at 0.5, 1, 3, 6, 12 h pooled (M203); macrophage cell line J774 + LPS +

IL-10 at 0.5, 1, 3, 5, 12 h pooled(M204); aerosol challenged mouse lung tissue, Th2 primers, aerosol OVA challenge 7, 14, 23 h pooled (see Garlisi, et al. (1995) Clinical Immunology and Immunopathology 75:75-83; X206); Nippostrongulus-infected lung tissue (see Coffman, et al. (1989) Science 245:308-310; X200); total adult lung, normal (O200); total lung, rag-1 (see Schwarz, et al. (1993) Immunodeficiency 4:249-252; O205); IL-10 K.O. spleen (see Kuhn, et al. (1991) Cell 75:263-274; X201); total adult spleen, normal (O201); total spleen, rag-1 (O207); IL-10 K.O. Peyer's patches (O202); total Peyer's patches, normal (O210); IL-10 K.O. mesenteric lymph nodes (X203); total mesenteric lymph nodes, normal (O211); IL-10 K.O. colon (X203); total colon, normal (O212); NOD mouse pancreas (see Makino, et al. (1980) Jikken Dobutsu 29:1-13; X205); total thymus, rag-1 (O208); total kidney, rag-1 (O209); total heart, rag-1 (O202); total brain, rag-1 (O203); total testes, rag-1 (O204); total liver, rag-1 (O206); rat normal joint tissue (O300); and rat arthritic joint tissue (X300).

The DTLR10 has been found to be highly expressed in precursor dendritic cell type 2 (pDC2). See, e.g., Rissoan, et al. (1999) Science 283:1183-1186; and Siegal, et al. (1999) Science 284:1835-1837. However, it is not expressed on monocytes. The restricted expression of DTLR10 reinforces the suggestions of a role for the receptor in host immune defense. The pDC2 cells are natural interferon producing cells (NIPC), which produce large amounts of IFNα in response to Herpes simplex virus infection.

V. Cloning of species counterparts of DTLRs

Various strategies are used to obtain species counterparts of these DTLRs, preferably from other primates. One method is by cross hybridization using closely related species DNA probes. It may be useful to go into evolutionarily similar species as intermediate

steps. Another method is by using specific PCR primers based on the identification of blocks of similarity or difference between particular species, e.g., human, genes, e.g., areas of highly conserved or nonconserved polypeptide or nucleotide sequence. Alternatively, antibodies may be used for expression cloning.

VI. Production of mammalian DTLR protein

An appropriate, e.g., GST, fusion construct is engineered for expression, e.g., in E. coli. For example, a mouse IGIF pGex plasmid is constructed and transformed into E. coli. Freshly transformed cells are grown in LB medium containing 50 µg/ml ampicillin and induced with IPTG (Sigma, St. Louis, MO). After overnight induction, the bacteria are harvested and the pellets containing the DTLR protein are isolated. The pellets are homogenized in TE buffer (50 mM Tris-base pH 8.0, 10 mM EDTA and 2 mM pefabloc) in 2 liters. This material is passed through a microfluidizer (Microfluidics, Newton, MA) three times. The fluidized supernatant is spun down on a Sorvall GS-3 rotor for 1 h at 13,000 rpm. The resulting supernatant containing the DTLR protein is filtered and passed over a glutathione-SEPHAROSE column equilibrated in 50 mM Tris-base pH 8.0. The fractions containing the DTLR-GST fusion protein are pooled and cleaved with thrombin (Enzyme Research Laboratories, Inc., South Bend, IN). The cleaved pool is then passed over a Q-SEPHAROSE column equilibrated in 50 mM Tris-base. Fractions containing DTLR are pooled and diluted in cold distilled H_2O, to lower the conductivity, and passed back over a fresh Q-Sepharose column, alone or in succession with an immunoaffinity antibody column.. Fractions containing the DTLR protein are pooled, aliquoted, and stored in the -70° C freezer.

Comparison of the CD spectrum with DTLR1 protein may suggest that the protein is correctly folded. See Hazuda, et al. (1969) J. Biol. Chem. 264:1689-1693.

VII. Biological Assays with DTLRs

Biological assays will generally be directed to the ligand binding feature of the protein or to the kinase/phosphatase activity of the receptor. The activity will typically be reversible, as are many other enzyme actions, and will mediate phosphatase or phosphorylase activities, which activities are easily measured by standard procedures. See, e.g., Hardie, et al. (eds. 1995) The Protein Kinase FactBook vols. I and II, Academic Press, San Diego, CA; Hanks, et al. (1991) Meth. Enzymol. 200:38-62; Hunter, et al. (1992) Cell 70:375-388; Lewin (1990) Cell 61:743-752; Pines, et al. (1991) Cold Spring Harbor Symp. Quant. Biol. 56:449-463; and Parker, et al. (1993) Nature 363:736-738.

The family of interleukin 1s contains molecules, each of which is an important mediator of inflammatory disease. For a comprehensive review, see Dinarello (1996) "Biologic basis for interleukin-1 in disease" Blood 87:2095-2147. There are suggestions that the various Toll ligands may play important roles in the initiation of disease, particularly inflammatory responses. The finding of novel proteins related to the IL-1 family furthers the identification of molecules that provide the molecular basis for initiation of disease and allow for the development of therapeutic strategies of increased range and efficacy.

VIII. Preparation of antibodies specific for, e.g., DTLR4

Inbred Balb/c mice are immunized intraperitoneally with recombinant forms of the protein, e.g., purified DTLR4 or stable transfected NIH-3T3 cells. Animals are boosted at appropriate time points with protein, with or without additional adjuvant, to further stimulate antibody production. Serum is collected, or hybridomas produced with harvested spleens.

Alternatively, Balb/c mice are immunized with cells transformed with the gene or fragments thereof, either endogenous or exogenous cells, or with isolated membranes enriched for expression of the antigen. Serum is collected at the appropriate time, typically after numerous further administrations. Various gene therapy techniques may be useful, e.g., in producing protein in situ, for generating an immune response.

Monoclonal antibodies may be made. For example, splenocytes are fused with an appropriate fusion partner and hybridomas are selected in growth medium by standard procedures. Hybridoma supernatants are screened for the presence of antibodies which bind to the desired DTLR, e.g., by ELISA or other assay. Antibodies which specifically recognize specific DTLR embodiments may also be selected or prepared.

In another method, synthetic peptides or purified protein are presented to an immune system to generate monoclonal or polyclonal antibodies. See, e.g., Coligan (1991) Current Protocols in Immunology Wiley/Greene; and Harlow and Lane (1989) Antibodies: A Laboratory Manual Cold Spring Harbor Press. In appropriate situations, the binding reagent is either labeled as described above, e.g., fluorescence or otherwise, or immobilized to a substrate for panning methods. Nucleic acids may also be introduced into cells in an animal to produce the antigen, which serves to elicit an immune response. See, e.g., Wang, et al. (1993) Proc. Nat'l. Acad. Sci. 90:4156-4160; Barry, et al. (1994) BioTechniques 16:616-619; and Xiang, et al. (1995) Immunity 2: 129-135.

IX. Production of fusion proteins with, e.g., DTLR5

Various fusion constructs are made with DTLR5. This portion of the gene is fused to an epitope tag, e.g., a FLAG tag, or to a two hybrid system construct. See, e.g., Fields and Song (1989) Nature 340:245-246.

The epitope tag may be used in an expression cloning procedure with detection with anti-FLAG antibodies to detect a binding partner, e.g., ligand for the respective DTLR5. The two hybrid system may also be used to isolate proteins which specifically bind to DTLR5.

X. Chromosomal mapping of DTLRs

Chromosome spreads are prepared. In situ hybridization is performed on chromosome preparations obtained from phytohemagglutinin-stimulated lymphocytes cultured for 72 h. 5-bromodeoxyuridine is added for the final seven hours of culture (60 µg/ml of medium), to ensure a posthybridization chromosomal banding of good quality.

An appropriate fragment, e.g., a PCR fragment, amplified with the help of primers on total B cell cDNA template, is cloned into an appropriate vector. The vector is labeled by nick-translation with ^{3}H. The radiolabeled probe is hybridized to metaphase spreads as described in Mattei, et al. (1985) Hum. Genet. 69:327-331.

After coating with nuclear track emulsion (KODAK NTB2), slides are exposed, e.g., for 18 days at 4° C. To avoid any slipping of silver grains during the banding procedure, chromosome spreads are first stained with buffered Giemsa solution and metaphase photographed. R-banding is then performed by the fluorochrome-photolysis-Giemsa (FPG) method and metaphases rephotographed before analysis.

Alternatively, FISH can be performed, as described above. The DTLR genes are located on different chromosomes. DTLR2 and DTLR3 are localized to human chromosome 4; DTLR4 is localized to human chromosome 9, and DTLR5 is localized to human chromosome 1. See Figures 4A-4D.

XI. Structure activity relationship

Information on the criticality of particular residues is determined using standard procedures and analysis. Standard mutagenesis analysis is performed, e.g., by generating many different variants at determined positions, e.g., at the positions identified above, and evaluating biological activities of the variants. This may be performed to the extent of determining positions which modify activity, or to focus on specific positions to determine the residues which can be substituted to either retain, block, or modulate biological activity.

Alternatively, analysis of natural variants can indicate what positions tolerate natural mutations. This may result from populational analysis of variation among individuals, or across strains or species. Samples from selected individuals are analyzed, e.g., by PCR analysis and sequencing. This allows evaluation of population polymorphisms.

XI. Isolation of a ligand for a DTLR

A DTLR can be used as a specific binding reagent to identify its binding partner, by taking advantage of its specificity of binding, much like an antibody would be used. A binding reagent is either labeled as described above, e.g., fluorescence or otherwise, or immobilized to a substrate for panning methods.

The binding composition is used to screen an expression library made from a cell line which expresses a binding partner, i.e., ligand, preferably membrane associated. Standard staining techniques are used to detect or sort surface expressed ligand, or surface expressing transformed cells are screened by panning. Screening of intracellular expression is performed by various staining or immunofluorescence procedures. See also McMahan, et al. (1991) EMBO J. 10:2821-2832.

For example, on day 0, precoat 2-chamber permanox slides with 1 ml per chamber of fibronectin, 10 ng/ml in PBS, for 30 min at room temperature. Rinse once with PBS. Then plate COS cells at 2-3 x 10^5 cells per chamber in 1.5 ml of growth media. Incubate overnight at 37° C.

On day 1 for each sample, prepare 0.5 ml of a solution of 66 µg/ml DEAE-dextran, 66 µM chloroquine, and 4 µg DNA in serum free DME. For each set, a positive control is prepared, e.g., of DTLR-FLAG cDNA at 1 and 1/200 dilution, and a negative mock. Rinse cells with serum free DME. Add the DNA solution and incubate 5 hr at 37° C. Remove the medium and add 0.5 ml 10% DMSO in DME for 2.5 min. Remove and wash once with DME. Add 1.5 ml growth medium and incubate overnight.

On day 2, change the medium. On days 3 or 4, the cells are fixed and stained. Rinse the cells twice with Hank's Buffered Saline Solution (HBSS) and fix in 4% paraformaldehyde (PFA)/glucose for 5 min. Wash 3X with HBSS. The slides may be stored at -80° C after all liquid is removed. For each chamber, 0.5 ml incubations are performed as follows. Add HBSS/saponin (0.1%) with 32 µl/ml of 1 M NaN_3 for 20 min. Cells are then washed with HBSS/saponin 1X. Add appropriate DTLR or DTLR/antibody complex to cells and incubate for 30 min. Wash cells twice with HBSS/saponin. If appropriate, add first antibody for 30 min. Add second antibody, e.g., Vector anti-mouse antibody, at 1/200 dilution, and incubate for 30 min. Prepare ELISA solution, e.g., Vector Elite ABC horseradish peroxidase solution, and preincubate for 30 min. Use, e.g., 1 drop of solution A (avidin) and 1 drop solution B (biotin) per 2.5 ml HBSS/saponin. Wash cells twice with HBSS/saponin. Add ABC HRP solution and incubate for 30 min. Wash cells twice with HBSS, second wash for 2 min, which closes cells. Then add Vector diaminobenzoic acid (DAB) for 5 to 10 min. Use 2 drops of buffer plus 4 drops DAB plus 2 drops of H_2O_2 per 5 ml of

glass distilled water. Carefully remove chamber and rinse slide in water. Air dry for a few minutes, then add 1 drop of Crystal Mount and a cover slip. Bake for 5 min at 85-90° C.

Evaluate positive staining of pools and progressively subclone to isolation of single genes responsible for the binding.

Alternatively, DTLR reagents are used to affinity purify or sort out cells expressing a putative ligand. See, e.g., Sambrook, et al. or Ausubel, et al.

Another strategy is to screen for a membrane bound receptor by panning. The receptor cDNA is constructed as described above. The ligand can be immobilized and used to immobilize expressing cells. Immobilization may be achieved by use of appropriate antibodies which recognize, e.g., a FLAG sequence of a DTLR fusion construct, or by use of antibodies raised against the first antibodies. Recursive cycles of selection and amplification lead to enrichment of appropriate clones and eventual isolation of receptor expressing clones.

Phage expression libraries can be screened by mammalian DTLRs. Appropriate label techniques, e.g., anti-FLAG antibodies, will allow specific labeling of appropriate clones.

All citations herein are incorporated herein by reference to the same extent as if each individual publication or patent application was specifically and individually indicated to be incorporated by reference.

Many modifications and variations of this invention can be made without departing from its spirit and scope, as will be apparent to those skilled in the art. The specific embodiments described herein are offered by way of example only, and the invention is to be limited by the terms of the appended claims, along with the full scope of equivalents to which such claims are entitled; and the

invention is not to be limited by the specific embodiments that have been presented herein by way of example.

Humans have two distinct types of dendritic cell (DC) precursors. Peripheral blood monocytes (pDC1) give rise to immature myeloid DCs after culturing with GMCSF and IL-4. These immature cells become mature myeloid DCs (DC1) after stimulation with CD40 ligand (CD40L). The CD4+CD3-CD11c- plasmacytoid cells (pDC2) from blood or tonsils give rise to a distinct type of immature DC after culture with IL-3, and differentiate into mature DCs (DC2) after CD40L stimulation. Rissoan, et al. (1999) Science 283:1183-1186.

Siegal, et al. (1999) Science 284:1835-1837, show that pDC2 is the "Natural Interferon Producing Cell" (IPC). Interferons (IFNs) are the most important cytokines in antiviral immune responses. "Natural IFN-producing cells" (NIPCs) in human blood express CD4 and major histocompatibility complex class II proteins, but have not been isolated and further characterized because of their rarity, rapid apoptosis, and lack of lineage markers. Purified NIPCs are here shown to be the CD4(+)CD11c- type 2 dendritic cell precursors (pDC2s), which produce 200 to 1000 times more IFN than other blood cells after microbial challenge. pDC2s are thus an effector cell type of the immune system, critical for antiviral and antitumor immune responses. They are implicated as important cells in HIV infected patients

Toll-like receptor (TLR) molecules belong to the IL-1/Toll receptor family. Ligands for TLR2 and TLR4 have been identified, and their functions are related to the host immune response to microbial antigen or injury. Takeuchi, et al. (1999) Immunity 11:443-451; and Noshino, et al. (1999) J. Immunol. 162:3749-3752. The pattern of expression of TLRs seem to be restricted. Muzio, et al. (2000) J. Immunol. 164:5998-6004. With these findings that: i) TLR10 is highly expressed and restricted in pDC2s, and ii) pDC2 is the NIPC, it is likely that TLR10 will play an important role in the host's innate immune response.

WHAT IS CLAIMED IS:

1. A composition of matter selected from the group consisting of:

a) a substantially pure or recombinant DTLR2 protein or peptide exhibiting identity over a length of at least about 12 amino acids to SEQ ID NO: 4;

b) a natural sequence DTLR2 of SEQ ID NO: 4;

c) a fusion protein comprising DTLR2 sequence;

d) a substantially pure or recombinant DTLR3 protein or peptide exhibiting identity over a length of at least about 12 amino acids to SEQ ID NO: 6;

e) a natural sequence DTLR3 of SEQ ID NO: 6;

f) a fusion protein comprising DTLR3 sequence;

g) a substantially pure or recombinant DTLR4 protein or peptide exhibiting identity over a length of at least about 12 amino acids to SEQ ID NO: 8;

h) a natural sequence DTLR4 of SEQ ID NO: 8;

i) a fusion protein comprising DTLR4 sequence;

j) a substantially pure or recombinant DTLR5 protein or peptide exhibiting identity over a length of at least about 12 amino acids to SEQ ID NO: 10;

k) a natural sequence DTLR5 comprising SEQ ID NO: 10;

l) a fusion protein comprising DTLR5 sequence;

m) a substantially pure or recombinant DTLR6 protein or peptide exhibiting identity over a length of at least about 12 amino acids to SEQ ID NO: 12, 28, or 30;

n) a natural sequence DTLR6 comprising SEQ ID NO: 12, 28, or 30;

o) a fusion protein comprising DTLR6 sequence;

p) a substantially pure or recombinant DTLR7 protein or peptide exhibiting identity over a length of at least about 12 amino acids to SEQ ID NO: 16, 18, or 37;

q) a natural sequence DTLR7 comprising SEQ ID NO: 16, 18, or 37;

r) a fusion protein comprising DTLR7 sequence;

s) a substantially pure or recombinant DTLR8 protein or peptide exhibiting identity over a length of at least about 12 amino acids to SEQ ID NO: 32 or 39;

t) a natural sequence DTLR8 comprising SEQ ID NO: 32 or 39;

u) a fusion protein comprising DTLR8 sequence;

v) a substantially pure or recombinant DTLR9 protein or peptide exhibiting identity over a length of at least about 12 amino acids to SEQ ID NO: 22 or 41;

w) a natural sequence DTLR9 comprising SEQ ID NO: 22 or 41;

x) a fusion protein comprising DTLR9 sequence;

y) a substantially pure or recombinant DTLR10 protein or peptide exhibiting identity over a length of at least about 12 amino acids to SEQ ID NO: 34, 43, or 45;

z) a natural sequence DTLR10 comprising SEQ ID NO: 34, 43, or 45;

zz) a fusion protein comprising DTLR10 sequence.

2. A substantially pure or isolated protein comprising a segment exhibiting sequence identity to a corresponding portion of a:

a) DTLR2 of Claim 1, and said identity is over at least:

a) about 15 amino acids;

b) about 19 amino acids; or

c) about 25 amino acids;

b) DTLR3 of Claim 1, and said identity is over at least:

a) about 15 amino acids;

b) about 19 amino acids; or
c) about 25 amino acids;

c) DTLR4 of Claim 1, and said identity is over at least:
a) about 15 amino acids;
b) about 19 amino acids; or
c) about 25 amino acids;

d) DTLR5 of Claim 1, and said identity is over at least:
a) about 15 amino acids;
b) about 19 amino acids; or
c) about 25 amino acids;

e) DTLR6 of Claim 1, and said identity is over at least:
a) about 15 amino acids;
b) about 19 amino acids; or
c) about 25 amino acids;

f) DTLR7 of Claim 1, and said identity is over at least:
a) about 15 amino acids;
b) about 19 amino acids; or
c) about 25 amino acids;

g) DTLR8 of Claim 1, and said identity is over at least:
a) about 15 amino acids;
b) about 19 amino acids; or
c) about 25 amino acids;

h) DTLR9 of Claim 1, and said identity is over at least:
a) about 15 amino acids;
b) about 19 amino acids; or
c) about 25 amino acids; or

i) DTLR10 of Claim 1, and said identity is over at least:
a) about 15 amino acids;
b) about 19 amino acids; or

c) about 25 amino acids.

3. The composition of matter of Claim 1, wherein said:

a) DTLR2:
   i) comprises a mature sequence of Table 2; or
   ii) lacks post-translational modification;
b) DTLR3:
   i) comprises a mature sequence of Table 3; or
   ii) lacks post-translational modification;
c) DTLR4:
   i) comprises a mature sequence of Table 4; or
   ii) lacks post-translational modification;
d) DTLR5:
   i) comprises a mature sequence of Table 5; or
   ii) lacks post-translational modification;
e) DTLR6:
   i) comprises a mature sequence of Table 6; or
   ii) lacks post-translational modification;
f) DTLR7:
   i) comprises a sequence of Table 7; or
   ii) lacks post-translational modification;
g) DTLR8:
   i) comprises a sequence of Table 8; or
   ii) lacks post-translational modification;
h) DTLR9:
   i) comprises a sequence of Table 9; or
   ii) lacks post-translational modification;
i) DTLR10:
   i) comprises a sequence of Table 10; or
   ii) lacks post-translational modification; or
j) protein or peptide:
   i) is from a warm blooded animal selected from a mammal, including a primate, such as a human;

ii) comprises at least one polypeptide segment of SEQ ID NO: 4, 6, 26, 10, 12, 28, 30, 16, 18, 37, 39, 32, 22, 34, 43, or 45;

iii) exhibits a plurality of said segments of identity;

iv) is a natural allelic variant of DTLR2, DTLR3, DTLR4, DTLR5, DTLR6, DTLR7, DTLR8, DTLR9, or DTLR10;

v) has a length at least about 30 amino acids;

vi) exhibits at least two non-overlapping epitopes which are specific for a primate DTLR2, DTLR3, DTLR4, DTLR5, DTLR6, DTLR7, DTLR8, DTLR9, or DTLR10;

vii) exhibits sequence identity over a length of at least about 35 amino acids to a primate DTLR2, DTLR3, DTLR4, DTLR5, DTLR6, DTLR7, DTLR8, DTLR9. or DTLR10;

viii) further exhibits at least two non-overlapping epitopes which are specific for a primate DTLR2, DTLR3, DTLR4, DTLR5, DTLR6, DTLR7, DTLR8, DTLR9, or DTLR10;

ix) exhibits identity over a length of at least about 20 amino acids to a rodent DTLR6;

x) is glycosylated;

xi) has a molecular weight of at least 100 kD with natural glycosylation;

xii) is a synthetic polypeptide;

xiii) is attached to a solid substrate;

xiv) is conjugated to another chemical moiety;

xv) is a 5-fold or less substitution from natural sequence; or

xvi) is a deletion or insertion variant from a natural sequence.

4. A composition comprising:

a) a sterile DTLR2 protein or peptide of Claim 1,

b) said DTLR2 protein or peptide of Claim 1 and a carrier, wherein said carrier is:
   i) an aqueous compound, including water, saline, and/or buffer; and/or
   ii) formulated for oral, rectal, nasal, topical, or parenteral administration;
c) a sterile DTLR3 protein or peptide of Claim 1;
d) said DTLR3 protein or peptide of Claim 1 and a carrier, wherein said carrier is:
   i) an aqueous compound, including water, saline, and/or buffer; and/or
   ii) formulated for oral, rectal, nasal, topical, or parenteral administration;
e) a sterile DTLR4 protein or peptide of Claim 1,
f) said DTLR4 protein or peptide of Claim 1 and a carrier, wherein said carrier is:
   i) an aqueous compound, including water, saline, and/or buffer; and/or
   ii) formulated for oral, rectal, nasal, topical, or parenteral administration;
g) a sterile DTLR5 protein or peptide of Claim 1;
h) said DTLR5 protein or peptide of Claim 1 and a carrier, wherein said carrier is:
   i) an aqueous compound, including water, saline, and/or buffer; and/or
   ii) formulated for oral, rectal, nasal, topical, or parenteral administration;
i) a sterile DTLR6 protein or peptide of Claim 1;
j) said DTLR6 protein or peptide of Claim 1 and a carrier, wherein said carrier is:
   i) an aqueous compound, including water, saline, and/or buffer; and/or
   ii) formulated for oral, rectal, nasal, topical, or parenteral administration;
k) a sterile DTLR7 protein or peptide of Claim 1;

l) said DTLR7 protein or peptide of Claim 1 and a carrier, wherein said carrier is:
   i) an aqueous compound, including water, saline, and/or buffer; and/or
   ii) formulated for oral, rectal, nasal, topical, or parenteral administration;
m) a sterile DTLR8 protein or peptide of Claim 1;
n) said DTLR8 protein or peptide of Claim 1 and a carrier, wherein said carrier is:
   i) an aqueous compound, including water, saline, and/or buffer; and/or
   ii) formulated for oral, rectal, nasal, topical, or parenteral administration;
o) a sterile DTLR9 protein or peptide of Claim 1;
p) said DTLR9 protein or peptide of Claim 1 and a carrier, wherein said carrier is:
   i) an aqueous compound, including water, saline, and/or buffer; and/or
   ii) formulated for oral, rectal, nasal, topical, or parenteral administration;
q) a sterile DTLR10 protein or peptide of Claim 1;
r) said DTLR10 protein or peptide of Claim 1 and a carrier, wherein said carrier is:
   i) an aqueous compound, including water, saline, and/or buffer; and/or
   ii) formulated for oral, rectal, nasal, topical, or parenteral administration;

5. The fusion protein of Claim 1, comprising:
a) mature protein comprising sequence of Table 2, 3, 4, 5, 6, 7, 8, 9, or 10;
b) a detection or purification tag, including a FLAG, His6, or Ig sequence; or
c) sequence of another receptor protein.

6. A kit comprising a protein or polypeptide of Claim 1, and:

a) a compartment comprising said protein or polypeptide; and/or

b) instructions for use or disposal of reagents in said kit.

7. A binding compound comprising an antigen binding site from an antibody, which specifically binds to a natural DTLR2, DTLR3, DTLR4, DTLR5, DTLR6, DTLR7, DTLR8, DTLR9, or DTLR10 protein of Claim 1, wherein:

a) said protein is a primate protein;

b) said binding compound is an Fv, Fab, or Fab2 fragment;

c) said binding compound is conjugated to another chemical moiety; or

d) said antibody:

i) is raised against a peptide sequence of a mature polypeptide of Table 2, 3, 4, 5, 6, 7, 8, 9, or 10;

ii) is raised against a mature DTLR2, DTLR3, DTLR4, DTLR5, DTLR6, DTLR7, DTLR8, DTLR9, or DTLR10;

iii) is raised to a purified human DTLR2, DTLR3, DTLR4, DTLR5, DTLR6, DTLR7, DTLR8, DTLR9, or DTLR10;

iv) is immunoselected;

v) is a polyclonal antibody;

vi) binds to a denatured DTLR2, DTLR3, DTLR4, DTLR5, DTLR6, DTLR7, DTLR8, DTLR9, or DTLR10;

vii) exhibits a Kd to antigen of at least 30 µM;

viii) is attached to a solid substrate, including a bead or plastic membrane;

ix) is in a sterile composition; or

x) is detectably labeled, including a radioactive or fluorescent label.

8. A kit comprising said binding compound of Claim 7, and:

a) a compartment comprising said binding compound; and/or

b) instructions for use or disposal of reagents in said kit.

9. A method of:

A) making an antibody of Claim 7, comprising immunizing an immune system with an immunogenic amount of:

a) a primate DTLR2;
b) a primate DTLR3;
c) a primate DTLR4;
d) a primate DTLR5;
e) a primate DTLR6;
f) a primate DTLR7;
g) a primate DTLR8;
h) a primate DTLR9; or
i) a primate DTLR10;

thereby causing said antibody to be produced; or

B) producing an antigen:antibody complex, comprising contacting an antibody of Claim 7 with:

a) a mammalian DTLR2 protein or peptide;
b) a mammalian DTLR3 protein or peptide;
c) a mammalian DTLR4 protein or peptide;
d) a mammalian DTLR5 protein or peptide;
e) a mammalian DTLR6 protein or peptide;
f) a mammalian DTLR7 protein or peptide;
g) a mammalian DTLR8 protein or peptide;
h) a mammalian DTLR9 protein or peptide; or
i) a mammalian DTLR10 protein or peptide;

thereby allowing said complex to form.

10. A composition comprising:

a) a sterile binding compound of Claim 7, or

b) said binding compound of Claim 7 and a carrier, wherein said carrier is:

i) an aqueous compound, including water, saline, and/or buffer; and/or

ii) formulated for oral, rectal, nasal, topical, or parenteral administration.

11. An isolated or recombinant nucleic acid encoding a protein or peptide or fusion protein of Claim 1, wherein:

a) said DTLR is from a mammal; or

b) said nucleic acid:

i) encodes an antigenic peptide sequence of Table 2, 3, 4, 5, 6, 7, 8, 9, or 10;

ii) encodes a plurality of antigenic peptide sequences of Table 2, 3, 4, 5, 6, 7, 8, 9, or 10;

iii) exhibits at least about 80% identity to a natural cDNA encoding said segment;

iv) is an expression vector;

v) further comprises an origin of replication;

vi) is from a natural source;

vii) comprises a detectable label;

viii) comprises synthetic nucleotide sequence;

ix) is less than 6 kb, preferably less than 3 kb;

x) is from a mammal, including a primate;

xi) comprises a natural full length coding sequence;

xii) is a hybridization probe for a gene encoding said DTLR;

xiii) comprises at least 17 contiguous nucleotides from Table 2, 3, 4, 5, 6, 7, 8, 9, or 10;

xiv) comprises at plurality of nonoverlapping segments of least 17 contiguous nucleotides from Table 2, 3, 4, 5, 6, 7, 8, 9, or 10; or

xv) is a PCR primer, PCR product, or mutagenesis primer.

12. A cell, tissue, or organ comprising a recombinant nucleic acid of Claim 11.

13. The cell of Claim 12, wherein said cell is:

a) a prokaryotic cell;
b) a eukaryotic cell;
c) a bacterial cell;
d) a yeast cell;
e) an insect cell;
f) a mammalian cell;
g) a mouse cell;
h) a primate cell; or
i) a human cell.

14. A kit comprising said nucleic acid of Claim 11, and:

a) a compartment comprising said nucleic acid;
b) a compartment further comprising a primate DTLR2, DTLR3, DTLR4, DTLR5, DTLR6, DTLR7, DTLR8, DTLR9, or DTLR10 protein or polypeptide; and/or
c) instructions for use or disposal of reagents in said kit.

15. A method of:

A) making a polypeptide, comprising expressing said nucleic acid of Claim 11, thereby producing said polypeptide; or

B) making a duplex nucleic acid, comprising contacting said nucleic acid of Claim 11 with a complementary nucleic acid, thereby allowing said duplex to form.

16. A nucleic acid which:

a) hybridizes under wash conditions of 30° C and less than 2M salt to SEQ ID NO: 3;

b) hybridizes under wash conditions of 30° C and less than 2 M salt to SEQ ID NO: 5;

c) hybridizes under wash conditions of 30° C and less than 2M salt to SEQ ID NO: 25;

d) hybridizes under wash conditions of 30° C and less than 2 M salt to SEQ ID NO: 9;

e) hybridizes under wash conditions of 30° C and less than 2M salt to SEQ ID NO: 11, 27, or 29;

f) hybridizes under wash conditions of 30° C and less than 2 M salt to SEQ ID NO: 15, 17, or 36;

g) hybridizes under wash conditions of 30° C and less than 2M salt to SEQ ID NO: 31 or 38;

h) hybridizes under wash conditions of 30° C and less than 2 M salt to SEQ ID NO: 21 or 40;

i) hybridizes under wash conditions of 30° C and less than 2 M salt to SEQ ID NO: 33, 35, 42, or 44;

j) exhibits at least about 85% identity over a stretch of at least about 30 nucleotides to a primate DTLR2;

k) exhibits at least about 85% identity over a stretch of at least about 30 nucleotides to a primate DTLR3;

l) exhibits at least about 85% identity over a stretch of at least about 30 nucleotides to a primate DTLR4;

m) exhibits at least about 85% identity over a stretch of at least about 30 nucleotides to a primate DTLR5;

n) exhibits at least about 85% identity over a stretch of at least about 30 nucleotides to a primate DTLR6;

o) exhibits at least about 85% identity over a stretch of at least about 30 nucleotides to a primate DTLR7;

p) exhibits at least about 85% identity over a stretch of at least about 30 nucleotides to a primate DTLR8;

q) exhibits at least about 85% identity over a stretch of at least about 30 nucleotides to a primate DTLR9; or

r) exhibits at least about 85% identity over a stretch of at least about 30 nucleotides to a primate DTLR10.

17. The nucleic acid of Claim 16, wherein:

a) said wash conditions are at 45° C and/or 500 mM salt; or

b) said identity is at least 90% and/or said stretch is at least 55 nucleotides.

18. The nucleic acid of Claim 17, wherein:

a) said wash conditions are at 55° C and/or 150 mM salt; or

b) said identity is at least 95% and/or said stretch is at least 75 nucleotides.

19. A method of producing a ligand:receptor complex, comprising contacting:

a) a substantially pure primate DTLR2, including a recombinant or synthetically produced protein, with candidate Toll ligand;

b) a substantially pure primate DTLR3, including a recombinant or synthetically produced protein, with candidate Toll ligand;

c) a substantially pure primate DTLR4, including a recombinant or synthetically produced protein, with candidate Toll ligand;

d) a substantially pure primate DTLR5, including a recombinant or synthetically produced protein, with candidate Toll ligand;

e) a substantially pure primate DTLR6, including a recombinant or synthetically produced protein, with candidate Toll ligand;

f) a substantially pure primate DTLR7, including a recombinant or synthetically produced protein, with candidate Toll ligand;

g) a substantially pure primate DTLR8, including a recombinant or synthetically produced protein, with candidate Toll ligand;

h) a substantially pure primate DTLR9, including a recombinant or synthetically produced protein, with candidate Toll ligand;

i) a substantially pure primate DTLR10, including a recombinant or synthetically produced protein, with candidate Toll ligand;

thereby allowing said complex to form.

20. A method of modulating physiology or development of a cell or tissue culture cells comprising contacting said cell with an agonist or antagonist of a mammalian DTLR2, DTLR3, DTLR4, DTLR5; DTLR6, DTLR7, DTLR8, DTLR9, or DTLR10.

21. The method of Claim 20, wherein said agonist or antagonist is of DTLR10, and said cell is a pDC2 cell.

SEQUENCE SUBMISSION

SEQ ID NO: 1 provides primate DTLR1 nucleotide sequence.
SEQ ID NO: 2 provides primate DTLR1 polypeptide sequence.
SEQ ID NO: 3 provides primate DTLR2 nucleotide sequence.
SEQ ID NO: 4 provides primate DTLR2 polypeptide sequence.
SEQ ID NO: 5 provides primate DTLR3 nucleotide sequence.
SEQ ID NO: 6 provides primate DTLR3 polypeptide sequence.
SEQ ID NO: 7 provides primate DTLR4 nucleotide sequence.
SEQ ID NO: 8 provides primate DTLR4 polypeptide sequence.
SEQ ID NO: 9 provides primate DTLR5 nucleotide sequence.
SEQ ID NO: 10 provides primate DTLR5 polypeptide sequence.
SEQ ID NO: 11 provides primate DTLR6 nucleotide sequence.
SEQ ID NO: 12 provides primate DTLR6 polypeptide sequence.
SEQ ID NO: 13 provides rodent DTLR6 nucleotide sequence.
SEQ ID NO: 14 provides rodent DTLR6 polypeptide sequence.
SEQ ID NO: 15 provides primate DTLR7 nucleotide sequence.
SEQ ID NO: 16 provides primate DTLR7 polypeptide sequence.
SEQ ID NO: 17 provides primate DTLR7 nucleotide sequence.
SEQ ID NO: 18 provides primate DTLR7 polypeptide sequence.
SEQ ID NO: 19 provides primate DTLR8 nucleotide sequence.
SEQ ID NO: 20 provides primate DTLR8 polypeptide sequence.
SEQ ID NO: 21 provides primate DTLR9 nucleotide sequence.
SEQ ID NO: 22 provides primate DTLR9 polypeptide sequence.
SEQ ID NO: 23 provides primate DTLR10 nucleotide sequence.
SEQ ID NO: 24 provides primate DTLR10 polypeptide sequence.
SEQ ID NO: 25 provides primate DTLR4 nucleotide sequence.
SEQ ID NO: 26 provides primate DTLR4 polypeptide sequence.
SEQ ID NO: 27 provides rodent DTLR6 nucleotide sequence.
SEQ ID NO: 28 provides rodent DTLR6 polypeptide sequence.
SEQ ID NO: 29 provides rodent DTLR6 nucleotide sequence.
SEQ ID NO: 30 provides rodent DTLR6 polypeptide sequence.
SEQ ID NO: 31 provides primate DTLR8 nucleotide sequence.
SEQ ID NO: 32 provides primate DTLR8 polypeptide sequence.
SEQ ID NO: 33 provides primate DTLR10 nucleotide sequence.
SEQ ID NO: 34 provides primate DTLR10 polypeptide sequence.
SEQ ID NO: 35 provides rodent DTLR10 nucleotide sequence.
SEQ ID NO: 36 provides primate DTLR7 nucleotide sequence.
SEQ ID NO: 37 provides primate DTLR7 polypeptide sequence.
SEQ ID NO: 38 provides primate DTLR8 nucleotide sequence.
SEQ ID NO: 39 provides primate DTLR8 polypeptide sequence.
SEQ ID NO: 40 provides primate DTLR9 nucleotide sequence.
SEQ ID NO: 41 provides primate DTLR9 polypeptide sequence.
SEQ ID NO: 42 provides primate DTLR10 nucleotide sequence.
SEQ ID NO: 43 provides primate DTLR10 polypeptide sequence.
SEQ ID NO: 44 provides rodent DTLR10 nucleotide sequence.
SEQ ID NO: 45 provides rodent DTLR10 polypeptide sequence.

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Gly Leu Ser Thr Thr His Leu Glu Lys Ser Ser Val Leu Pro Ile Ala
        125                 130                 135

cat ttg aat atc agc aag gtc ttg ctg gtc tta gga gag act tat ggg   528
His Leu Asn Ile Ser Lys Val Leu Leu Val Leu Gly Glu Thr Tyr Gly
    140                 145                 150

gaa aaa gaa gac cct gag ggc ctt caa gac ttt aac act gag agt ctg   576
Glu Lys Glu Asp Pro Glu Gly Leu Gln Asp Phe Asn Thr Glu Ser Leu
155                 160                 165                 170

cac att gtg ttc ccc aca aac aaa gaa ttc cat ttt att ttg gat gtg   624
His Ile Val Phe Pro Thr Asn Lys Glu Phe His Phe Ile Leu Asp Val
                175                 180                 185

tca gtc aag act gta gca aat ctg gaa cta tct aat atc aaa tgt gtg   672
Ser Val Lys Thr Val Ala Asn Leu Glu Leu Ser Asn Ile Lys Cys Val
            190                 195                 200

cta gaa gat aac aaa tgt tct tac ttc cta agt att ctg gcg aaa ctt   720
Leu Glu Asp Asn Lys Cys Ser Tyr Phe Leu Ser Ile Leu Ala Lys Leu
        205                 210                 215

caa aca aat cca aag tta tca agt ctt acc tta aac aac att gaa aca   768
Gln Thr Asn Pro Lys Leu Ser Ser Leu Thr Leu Asn Asn Ile Glu Thr
    220                 225                 230

act tgg aat tct ttc att agg atc ctc caa cta gtt tgg cat aca act   816
Thr Trp Asn Ser Phe Ile Arg Ile Leu Gln Leu Val Trp His Thr Thr
235                 240                 245                 250

gta tgg tat ttc tca att tca aac gtg aag cta cag ggt cag ctg gac   864
Val Trp Tyr Phe Ser Ile Ser Asn Val Lys Leu Gln Gly Gln Leu Asp
                255                 260                 265

ttc aga gat ttt gat tat tct ggc act tcc ttg aag gcc ttg tct ata   912
Phe Arg Asp Phe Asp Tyr Ser Gly Thr Ser Leu Lys Ala Leu Ser Ile
            270                 275                 280

cac caa gtt gtc agc gat gtg ttc ggt ttt ccg caa agt tat atc tat   960
His Gln Val Val Ser Asp Val Phe Gly Phe Pro Gln Ser Tyr Ile Tyr
        285                 290                 295

gaa atc ttt tcg aat atg aac atc aaa aat ttc aca gtg tct ggt aca  1008
Glu Ile Phe Ser Asn Met Asn Ile Lys Asn Phe Thr Val Ser Gly Thr
    300                 305                 310

cgc atg gtc cac atg ctt tgc cca tcc aaa att agc ccg ttc ctg cat  1056
Arg Met Val His Met Leu Cys Pro Ser Lys Ile Ser Pro Phe Leu His
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     315                 320                 325                 330

     ttg gat ttt tcc aat aat ctc tta aca gac acg gtt ttt gaa aat tgt   1104
     Leu Asp Phe Ser Asn Asn Leu Leu Thr Asp Thr Val Phe Glu Asn Cys
 5                   335                 340                 345

     ggg cac ctt act gag ttg gag aca ctt att tta caa atg aat caa tta   1152
     Gly His Leu Thr Glu Leu Glu Thr Leu Ile Leu Gln Met Asn Gln Leu
                 350                 355                 360
10
     aaa gaa ctt tca aaa ata gct gaa atg act aca cag atg aag tct ctg   1200
     Lys Glu Leu Ser Lys Ile Ala Glu Met Thr Thr Gln Met Lys Ser Leu
             365                 370                 375

15   caa caa ttg gat att agc cag aat tct gta agc tat gat gaa aag aaa   1248
     Gln Gln Leu Asp Ile Ser Gln Asn Ser Val Ser Tyr Asp Glu Lys Lys
         380                 385                 390

     gga gac tgt tct tgg act aaa agt tta tta agt tta aat atg tct tca   1296
20   Gly Asp Cys Ser Trp Thr Lys Ser Leu Leu Ser Leu Asn Met Ser Ser
     395                 400                 405                 410

     aat ata ctt act gac act att ttc aga tgt tta cct ccc agg atc aag   1344
     Asn Ile Leu Thr Asp Thr Ile Phe Arg Cys Leu Pro Pro Arg Ile Lys
25                   415                 420                 425

     gta ctt gat ctt cac agc aat aaa ata aag agc att cct aaa caa gtc   1392
     Val Leu Asp Leu His Ser Asn Lys Ile Lys Ser Ile Pro Lys Gln Val
                 430                 435                 440
30
     gta aaa ctg gaa gct ttg caa gaa ctc aat gtt gct ttc aat tct tta   1440
     Val Lys Leu Glu Ala Leu Gln Glu Leu Asn Val Ala Phe Asn Ser Leu
             445                 450                 455

35   act gac ctt cct gga tgt ggc agc ttt agc agc ctt tct gta ttg atc   1488
     Thr Asp Leu Pro Gly Cys Gly Ser Phe Ser Ser Leu Ser Val Leu Ile
         460                 465                 470

     att gat cac aat tca gtt tcc cac cca tca gct gat ttc ttc cag agc   1536
40   Ile Asp His Asn Ser Val Ser His Pro Ser Ala Asp Phe Phe Gln Ser
     475                 480                 485                 490

     tgc cag aag atg agg tca ata aaa gca ggg gac aat cca ttc caa tgt   1584
     Cys Gln Lys Met Arg Ser Ile Lys Ala Gly Asp Asn Pro Phe Gln Cys
45                   495                 500                 505

     acc tgt gag ctc gga gaa ttt gtc aaa aat ata gac caa gta tca agt   1632
     Thr Cys Glu Leu Gly Glu Phe Val Lys Asn Ile Asp Gln Val Ser Ser
                 510                 515                 520
50
     gaa gtg tta gag ggc tgg cct gat tct tat aag tgt gac tac ccg gaa   1680
     Glu Val Leu Glu Gly Trp Pro Asp Ser Tyr Lys Cys Asp Tyr Pro Glu
             525                 530                 535

55   agt tat aga gga acc cta cta aag gac ttt cac atg tct gaa tta tcc   1728
     Ser Tyr Arg Gly Thr Leu Leu Lys Asp Phe His Met Ser Glu Leu Ser
         540                 545                 550
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tgc aac ata act ctg ctg atc gtc acc atc gtt gcc acc atg ctg gtg   1776
Cys Asn Ile Thr Leu Leu Ile Val Thr Ile Val Ala Thr Met Leu Val
555                 560                 565                 570

ttg gct gtg act gtg acc tcc ctc tgc atc tac ttg gat ctg ccc tgg   1824
Leu Ala Val Thr Val Thr Ser Leu Cys Ile Tyr Leu Asp Leu Pro Trp
                575                 580                 585

tat ctc agg atg gtg tgc cag tgg acc cag acc cgg cgc agg gcc agg   1872
Tyr Leu Arg Met Val Cys Gln Trp Thr Gln Thr Arg Arg Arg Ala Arg
            590                 595                 600

aac ata ccc tta gaa gaa ctc caa aga aat ctc cag ttt cat gca ttt   1920
Asn Ile Pro Leu Glu Glu Leu Gln Arg Asn Leu Gln Phe His Ala Phe
        605                 610                 615

att tca tat agt ggg cac gat tct ttc tgg gtg aag aat gaa tta ttg   1968
Ile Ser Tyr Ser Gly His Asp Ser Phe Trp Val Lys Asn Glu Leu Leu
    620                 625                 630

cca aac cta gag aaa gaa ggt atg cag att tgc ctt cat gag aga aac   2016
Pro Asn Leu Glu Lys Glu Gly Met Gln Ile Cys Leu His Glu Arg Asn
635                 640                 645                 650

ttt gtt cct ggc aag agc att ctg gaa aat atc atc acc tgc att gag   2064
Phe Val Pro Gly Lys Ser Ile Val Glu Asn Ile Ile Thr Cys Ile Glu
                655                 660                 665

aag agt tac aag tcc atc ttt gtt ttg tct ccc aac ttt gtc cag agt   2112
Lys Ser Tyr Lys Ser Ile Phe Val Leu Ser Pro Asn Phe Val Gln Ser
            670                 675                 680

gaa tgg tgc cat tat gaa ctc tac ttt gcc cat cac aat ctc ttt cat   2160
Glu Trp Cys His Tyr Glu Leu Tyr Phe Ala His His Asn Leu Phe His
        685                 690                 695

gaa gga tct aat agc tta atc ctg atc ttg ctg gaa ccc att ccg cag   2208
Glu Gly Ser Asn Ser Leu Ile Leu Ile Leu Leu Glu Pro Ile Pro Gln
    700                 705                 710

tac tcc att cct agc agt tat cac aag ctc aaa agt ctc atg gcc agg   2256
Tyr Ser Ile Pro Ser Ser Tyr His Lys Leu Lys Ser Leu Met Ala Arg
715                 720                 725                 730

agg act tat ttg gaa tgg ccc aag gaa aag agc aaa cgt ggc ctt ttt   2304
Arg Thr Tyr Leu Glu Trp Pro Lys Glu Lys Ser Lys Arg Gly Leu Phe
                735                 740                 745

tgg gct aac tta agg gca gcc att aat att aag ctg aca gag caa gca   2352
Trp Ala Asn Leu Arg Ala Ala Ile Asn Ile Lys Leu Thr Glu Gln Ala
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Lys Lys
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-5              -1  1                   5                  10

Lys Asn Gly Leu Ile His Val Pro Lys Asp Leu Ser Gln Lys Thr Thr
            15                  20                  25

Ile Leu Asn Ile Ser Gln Asn Tyr Ile Ser Glu Leu Trp Thr Ser Asp
        30                  35                  40

Ile Leu Ser Leu Ser Lys Leu Arg Ile Leu Ile Ile Ser His Asn Arg
    45                  50                  55

Ile Gln Tyr Leu Asp Ile Ser Val Phe Lys Phe Asn Gln Glu Leu Glu
60                  65                  70

Tyr Leu Asp Leu Ser His Asn Lys Leu Val Lys Ile Ser Cys His Pro
75                  80                  85                  90

Thr Val Asn Leu Lys His Leu Asp Leu Ser Phe Asn Ala Phe Asp Ala
                95                  100                 105

Leu Pro Ile Cys Lys Glu Phe Gly Asn Met Ser Gln Leu Lys Phe Leu
            110                 115                 120

Gly Leu Ser Thr Thr His Leu Glu Lys Ser Ser Val Leu Pro Ile Ala
        125                 130                 135

His Leu Asn Ile Ser Lys Val Leu Leu Val Leu Gly Glu Thr Tyr Gly
    140                 145                 150

Glu Lys Glu Asp Pro Glu Gly Leu Gln Asp Phe Asn Thr Glu Ser Leu
155                 160                 165                 170

His Ile Val Phe Pro Thr Asn Lys Glu Phe His Phe Ile Leu Asp Val
                175                 180                 185

Ser Val Lys Thr Val Ala Asn Leu Glu Leu Ser Asn Ile Lys Cys Val
            190                 195                 200

Leu Glu Asp Asn Lys Cys Ser Tyr Phe Leu Ser Ile Leu Ala Lys Leu
        205                 210                 215

Gln Thr Asn Pro Lys Leu Ser Ser Leu Thr Leu Asn Asn Ile Glu Thr
    220                 225                 230

Thr Trp Asn Ser Phe Ile Arg Ile Leu Gln Leu Val Trp His Thr Thr
235                 240                 245                 250

Val Trp Tyr Phe Ser Ile Ser Asn Val Lys Leu Gln Gly Gln Leu Asp
                255                 260                 265
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Phe Arg Asp Phe Asp Tyr Ser Gly Thr Ser Leu Lys Ala Leu Ser Ile
270 275 280

His Gln Val Val Ser Asp Val Phe Gly Phe Pro Gln Ser Tyr Ile Tyr
285 290 295

Glu Ile Phe Ser Asn Met Asn Ile Lys Asn Phe Thr Val Ser Gly Thr
300 305 310

Arg Met Val His Met Leu Cys Pro Ser Lys Ile Ser Pro Phe Leu His
315 320 325 330

Leu Asp Phe Ser Asn Asn Leu Leu Thr Asp Thr Val Phe Glu Asn Cys
335 340 345

Gly His Leu Thr Glu Leu Glu Thr Leu Ile Leu Gln Met Asn Gln Leu
350 355 360

Lys Glu Leu Ser Lys Ile Ala Glu Met Thr Thr Gln Met Lys Ser Leu
365 370 375

Gln Gln Leu Asp Ile Ser Gln Asn Ser Val Ser Tyr Asp Glu Lys Lys
380 385 390

Gly Asp Cys Ser Trp Thr Lys Ser Leu Leu Ser Leu Asn Met Ser Ser
395 400 405 410

Asn Ile Leu Thr Asp Thr Ile Phe Arg Cys Leu Pro Pro Arg Ile Lys
415 420 425

Val Leu Asp Leu His Ser Asn Lys Ile Lys Ser Ile Pro Lys Gln Val
430 435 440

Val Lys Leu Glu Ala Leu Gln Glu Leu Asn Val Ala Phe Asn Ser Leu
445 450 455

Thr Asp Leu Pro Gly Cys Gly Ser Phe Ser Ser Leu Ser Val Leu Ile
460 465 470

Ile Asp His Asn Ser Val Ser His Pro Ser Ala Asp Phe Phe Gln Ser
475 480 485 490

Cys Gln Lys Met Arg Ser Ile Lys Ala Gly Asp Asn Pro Phe Gln Cys
495 500 505

Thr Cys Glu Leu Gly Glu Phe Val Lys Asn Ile Asp Gln Val Ser Ser
510 515 520

Glu Val Leu Glu Gly Trp Pro Asp Ser Tyr Lys Cys Asp Tyr Pro Glu
525 530 535

Ser Tyr Arg Gly Thr Leu Leu Lys Asp Phe His Met Ser Glu Leu Ser
540 545 550

Cys Asn Ile Thr Leu Leu Ile Val Thr Ile Val Ala Thr Met Leu Val
555 560 565 570

Leu Ala Val Thr Val Thr Ser Leu Cys Ile Tyr Leu Asp Leu Pro Trp

575 580 585

Tyr Leu Arg Met Val Cys Gln Trp Thr Gln Thr Arg Arg Arg Ala Arg
590 595 600

Asn Ile Pro Leu Glu Glu Leu Gln Arg Asn Leu Gln Phe His Ala Phe
605 610 615

Ile Ser Tyr Ser Gly His Asp Ser Phe Trp Val Lys Asn Glu Leu Leu
620 625 630

Pro Asn Leu Glu Lys Glu Gly Met Gln Ile Cys Leu His Glu Arg Asn
635 640 645 650

Phe Val Pro Gly Lys Ser Ile Val Glu Asn Ile Ile Thr Cys Ile Glu
655 660 665

Lys Ser Tyr Lys Ser Ile Phe Val Leu Ser Pro Asn Phe Val Gln Ser
670 675 680

Glu Trp Cys His Tyr Glu Leu Tyr Phe Ala His His Asn Leu Phe His
685 690 695

Glu Gly Ser Asn Ser Leu Ile Leu Ile Leu Leu Glu Pro Ile Pro Gln
700 705 710

Tyr Ser Ile Pro Ser Ser Tyr His Lys Leu Lys Ser Leu Met Ala Arg
715 720 725 730

Arg Thr Tyr Leu Glu Trp Pro Lys Glu Lys Ser Lys Arg Gly Leu Phe
735 740 745

Trp Ala Asn Leu Arg Ala Ala Ile Asn Ile Lys Leu Thr Glu Gln Ala
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Lys Lys

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ctc tcc aag gaa gaa tcc tcc aat cag gct tct ctg tct tgt gac cgc   96
Leu Ser Lys Glu Glu Ser Ser Asn Gln Ala Ser Leu Ser Cys Asp Arg
     -5              -1  1                   5                  10

aat ggt atc tgc aag ggc agc tca gga tct tta aac tcc att ccc tca   144
Asn Gly Ile Cys Lys Gly Ser Ser Gly Ser Leu Asn Ser Ile Pro Ser
                15                  20                  25

ggg ctc aca gaa gct gta aaa agc ctt gac ctg tcc aac aac agg atc   192
Gly Leu Thr Glu Ala Val Lys Ser Leu Asp Leu Ser Asn Asn Arg Ile
            30                  35                  40

acc tac att agc aac agt gac cta cag agg tgt gtg aac ctc cag gct   240
Thr Tyr Ile Ser Asn Ser Asp Leu Gln Arg Cys Val Asn Leu Gln Ala
        45                  50                  55

ctg gtg ctg aca tcc aat gga att aac aca ata gag gaa gat tct ttt   288
Leu Val Leu Thr Ser Asn Gly Ile Asn Thr Ile Glu Glu Asp Ser Phe
    60                  65                  70

tct tcc ctg ggc agt ctt gaa cat tta gac tta tcc tat aat tac tta   336
Ser Ser Leu Gly Ser Leu Glu His Leu Asp Leu Ser Tyr Asn Tyr Leu
75                  80                  85                  90

tct aat tta tcg tct tcc tgg ttc aag ccc ctt tct tct tta aca ttc   384
Ser Asn Leu Ser Ser Ser Trp Phe Lys Pro Leu Ser Ser Leu Thr Phe
                95                  100                 105

tta aac tta ctg gga aat cct tac aaa acc cta ggg gaa aca tct ctt   432
Leu Asn Leu Leu Gly Asn Pro Tyr Lys Thr Leu Gly Glu Thr Ser Leu
            110                 115                 120

ttt tct cat ctc aca aaa ttg caa atc ctg aga gtg gga aat atg gac   480
Phe Ser His Leu Thr Lys Leu Gln Ile Leu Arg Val Gly Asn Met Asp
        125                 130                 135

acc ttc act aag att caa aga aaa gat ttt gct gga ctt acc ttc ctt   528
Thr Phe Thr Lys Ile Gln Arg Lys Asp Phe Ala Gly Leu Thr Phe Leu
    140                 145                 150

gag gaa ctt gag att gat gct tca gat cta cag agc tat gag cca aaa   576
Glu Glu Leu Glu Ile Asp Ala Ser Asp Leu Gln Ser Tyr Glu Pro Lys
155                 160                 165                 170

agt ttg aag tca att cag aac gta agt cat ctg atc ctt cat atg aag   624
Ser Leu Lys Ser Ile Gln Asn Val Ser His Leu Ile Leu His Met Lys
                175                 180                 185

cag cat att tta ctg ctg gag att ttt gta gat gtt aca agt tcc gtg   672
Gln His Ile Leu Leu Leu Glu Ile Phe Val Asp Val Thr Ser Ser Val
            190                 195                 200

gaa tgt ttg gaa ctg cga gat act gat ttg gac act ttc cat ttt tca   720
Glu Cys Leu Glu Leu Arg Asp Thr Asp Leu Asp Thr Phe His Phe Ser
        205                 210                 215
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gaa cta tcc act ggt gaa aca aat tca ttg att aaa aag ttt aca ttt   768
Glu Leu Ser Thr Gly Glu Thr Asn Ser Leu Ile Lys Lys Phe Thr Phe
    220                 225                 230

aga aat gtg aaa atc acc gat gaa agt ttg ttt cag gtt atg aaa ctt   816
Arg Asn Val Lys Ile Thr Asp Glu Ser Leu Phe Gln Val Met Lys Leu
235                 240                 245                 250

ttg aat cag att tct gga ttg tta gaa tta gag ttt gat gac tgt acc   864
Leu Asn Gln Ile Ser Gly Leu Leu Glu Leu Glu Phe Asp Asp Cys Thr
                255                 260                 265

ctt aat gga gtt ggt aat ttt aga gca tct gat aat gac aga gtt ata   912
Leu Asn Gly Val Gly Asn Phe Arg Ala Ser Asp Asn Asp Arg Val Ile
            270                 275                 280

gat cca ggt aaa gtg gaa acg tta aca atc cgg agg ctg cat att cca   960
Asp Pro Gly Lys Val Glu Thr Leu Thr Ile Arg Arg Leu His Ile Pro
        285                 290                 295

agg ttt tac tta ttt tat gat ctg agc act tta tat tca ctt aca gaa  1008
Arg Phe Tyr Leu Phe Tyr Asp Leu Ser Thr Leu Tyr Ser Leu Thr Glu
    300                 305                 310

aga gtt aaa aga atc aca gta gaa aac agt aaa gtt ttt ctg gtt cct  1056
Arg Val Lys Arg Ile Thr Val Glu Asn Ser Lys Val Phe Leu Val Pro
315                 320                 325                 330

tgt tta ctt tca caa cat tta aaa tca tta gaa tac ttg gat ctc agt  1104
Cys Leu Leu Ser Gln His Leu Lys Ser Leu Glu Tyr Leu Asp Leu Ser
                335                 340                 345

gaa aat ttg atg gtt gaa gaa tac ttg aaa aat tca gcc tgt gag gat  1152
Glu Asn Leu Met Val Glu Glu Tyr Leu Lys Asn Ser Ala Cys Glu Asp
            350                 355                 360

gcc tgg ccc tct cta caa act tta att tta agg caa aat cat ttg gca  1200
Ala Trp Pro Ser Leu Gln Thr Leu Ile Leu Arg Gln Asn His Leu Ala
        365                 370                 375

tca ttg gaa aaa acc gga gag act ttg ctc act ctg aaa aac ttg act  1248
Ser Leu Glu Lys Thr Gly Glu Thr Leu Leu Thr Leu Lys Asn Leu Thr
    380                 385                 390

aac att gat atc agt aag aat agt ttt cat tct atg cct gaa act tgt  1296
Asn Ile Asp Ile Ser Lys Asn Ser Phe His Ser Met Pro Glu Thr Cys
395                 400                 405                 410

cag tgg cca gaa aag atg aaa tat ttg aac tta tcc agc aca cga ata  1344
Gln Trp Pro Glu Lys Met Lys Tyr Leu Asn Leu Ser Ser Thr Arg Ile
                415                 420                 425

cac agt gta aca ggc tgc att ccc aag aca ctg gaa att tta gat gtt  1392
His Ser Val Thr Gly Cys Ile Pro Lys Thr Leu Glu Ile Leu Asp Val
            430                 435                 440

agc aac aac aat ctc aat tta ttt tct ttg aat ttg ccg caa ctc aaa  1440
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Ser Asn Asn Asn Leu Asn Leu Phe Ser Leu Asn Leu Pro Gln Leu Lys
        445                 450                 455

gaa ctt tat att tcc aga aat aag ttg atg act cta cca gat gcc tcc  1488
Glu Leu Tyr Ile Ser Arg Asn Lys Leu Met Thr Leu Pro Asp Ala Ser
    460                 465                 470

ctc tta ccc atg tta cta gta ttg aaa atc agt agg aat gca ata act  1536
Leu Leu Pro Met Leu Leu Val Leu Lys Ile Ser Arg Asn Ala Ile Thr
475                 480                 485                 490

acg ttt tct aag gag caa ctt gac tca ttt cac aca ctg aag act ttg  1584
Thr Phe Ser Lys Glu Gln Leu Asp Ser Phe His Thr Leu Lys Thr Leu
                495                 500                 505

gaa gct ggt ggc aat aac ttc att tgc tcc tgt gaa ttc ctc tcc ttc  1632
Glu Ala Gly Gly Asn Asn Phe Ile Cys Ser Cys Glu Phe Leu Ser Phe
            510                 515                 520

act cag gag cag caa gca ctg gcc aaa gtc ttg att gat tgg cca gca  1680
Thr Gln Glu Gln Gln Ala Leu Ala Lys Val Leu Ile Asp Trp Pro Ala
        525                 530                 535

aat tac ctg tgt gac tct cca tcc cat gtg cgt ggc cag cag gtt cag  1728
Asn Tyr Leu Cys Asp Ser Pro Ser His Val Arg Gly Gln Gln Val Gln
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gat gtc cgc ctc tcg gtg tcg gaa tgt cac agg aca gca ctg gtg tct  1776
Asp Val Arg Leu Ser Val Ser Glu Cys His Arg Thr Ala Leu Val Ser
555                 560                 565                 570

ggc atg tgc tgt gct ctg ttc ctg ctg atc ctg ctc acg ggg gtc ctg  1824
Gly Met Cys Cys Ala Leu Phe Leu Leu Ile Leu Leu Thr Gly Val Leu
                575                 580                 585

tgc cac cgt ttc cat ggc ctg tgg tat atg aaa atg atg tgg gcc tgg  1872
Cys His Arg Phe His Gly Leu Trp Tyr Met Lys Met Met Trp Ala Trp
            590                 595                 600

ctc cag gcc aaa agg aag ccc agg aaa gct ccc agc agg aac atc tgc  1920
Leu Gln Ala Lys Arg Lys Pro Arg Lys Ala Pro Ser Arg Asn Ile Cys
        605                 610                 615

tat gat gca ttt gtt tct tac agt gag cgg gat gcc tac tgg gtg gag  1968
Tyr Asp Ala Phe Val Ser Tyr Ser Glu Arg Asp Ala Tyr Trp Val Glu
    620                 625                 630

aac ctt atg gtc cag gag ctg gag aac ttc aat ccc ccc ttc aag ttg  2016
Asn Leu Met Val Gln Glu Leu Glu Asn Phe Asn Pro Pro Phe Lys Leu
635                 640                 645                 650

tgt ctt cat aag cgg gac ttc att cct ggc aag tgg atc att gac aat  2064
Cys Leu His Lys Arg Asp Phe Ile Pro Gly Lys Trp Ile Ile Asp Asn
                655                 660                 665

atc att gac tcc att gaa aag agc cac aaa act gtc ttt gtg ctt tct  2112
Ile Ile Asp Ser Ile Glu Lys Ser His Lys Thr Val Phe Val Leu Ser
            670                 675                 680
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gaa aac ttt gtg aag agt gag tgg tgc aag tat gaa ctg gac ttc tcc   2160
Glu Asn Phe Val Lys Ser Glu Trp Cys Lys Tyr Glu Leu Asp Phe Ser
        685                 690                 695

cat ttc cgt ctt ttt gaa gag aac aat gat gct gcc att ctc att ctt   2208
His Phe Arg Leu Phe Glu Glu Asn Asn Asp Ala Ala Ile Leu Ile Leu
    700                 705                 710

ctg gag ccc att gag aaa aaa gcc att ccc cag cgc ttc tgc aag ctg   2256
Leu Glu Pro Ile Glu Lys Lys Ala Ile Pro Gln Arg Phe Cys Lys Leu
715                 720                 725                 730

cgg aag ata atg aac acc aag acc tac ctg gag tgg ccc atg gac gag   2304
Arg Lys Ile Met Asn Thr Lys Thr Tyr Leu Glu Trp Pro Met Asp Glu
                735                 740                 745

gct cag cgg gaa gga ttt tgg gta aat ctg aga gct gcg ata aag tcc   2352
Ala Gln Arg Glu Gly Phe Trp Val Asn Leu Arg Ala Ala Ile Lys Ser
            750                 755                 760

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Asn Gly Ile Cys Lys Gly Ser Ser Gly Ser Leu Asn Ser Ile Pro Ser
                15                  20                  25

Gly Leu Thr Glu Ala Val Lys Ser Leu Asp Leu Ser Asn Asn Arg Ile
            30                  35                  40

Thr Tyr Ile Ser Asn Ser Asp Leu Gln Arg Cys Val Asn Leu Gln Ala
        45                  50                  55

Leu Val Leu Thr Ser Asn Gly Ile Asn Thr Ile Glu Glu Asp Ser Phe
    60                  65                  70

Ser Ser Leu Gly Ser Leu Glu His Leu Asp Leu Ser Tyr Asn Tyr Leu
75                  80                  85                  90

Ser Asn Leu Ser Ser Ser Trp Phe Lys Pro Leu Ser Ser Leu Thr Phe
                95                  100                 105

Leu Asn Leu Leu Gly Asn Pro Tyr Lys Thr Leu Gly Glu Thr Ser Leu
            110                 115                 120

Phe Ser His Leu Thr Lys Leu Gln Ile Leu Arg Val Gly Asn Met Asp
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                125                 130                 135

Thr Phe Thr Lys Ile Gln Arg Lys Asp Phe Ala Gly Leu Thr Phe Leu
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Glu Glu Leu Glu Ile Asp Ala Ser Asp Leu Gln Ser Tyr Glu Pro Lys
155                 160                 165                 170

Ser Leu Lys Ser Ile Gln Asn Val Ser His Leu Ile Leu His Met Lys
                175                 180                 185

Gln His Ile Leu Leu Leu Glu Ile Phe Val Asp Val Thr Ser Ser Val
            190                 195                 200

Glu Cys Leu Glu Leu Arg Asp Thr Asp Leu Asp Thr Phe His Phe Ser
        205                 210                 215

Glu Leu Ser Thr Gly Glu Thr Asn Ser Leu Ile Lys Lys Phe Thr Phe
    220                 225                 230

Arg Asn Val Lys Ile Thr Asp Glu Ser Leu Phe Gln Val Met Lys Leu
235                 240                 245                 250

Leu Asn Gln Ile Ser Gly Leu Leu Glu Leu Glu Phe Asp Asp Cys Thr
                255                 260                 265

Leu Asn Gly Val Gly Asn Phe Arg Ala Ser Asp Asn Asp Arg Val Ile
            270                 275                 280

Asp Pro Gly Lys Val Glu Thr Leu Thr Ile Arg Arg Leu His Ile Pro
        285                 290                 295

Arg Phe Tyr Leu Phe Tyr Asp Leu Ser Thr Leu Tyr Ser Leu Thr Glu
    300                 305                 310

Arg Val Lys Arg Ile Thr Val Glu Asn Ser Lys Val Phe Leu Val Pro
315                 320                 325                 330

Cys Leu Leu Ser Gln His Leu Lys Ser Leu Glu Tyr Leu Asp Leu Ser
                335                 340                 345

Glu Asn Leu Met Val Glu Glu Tyr Leu Lys Asn Ser Ala Cys Glu Asp
            350                 355                 360

Ala Trp Pro Ser Leu Gln Thr Leu Ile Leu Arg Gln Asn His Leu Ala
        365                 370                 375

Ser Leu Glu Lys Thr Gly Glu Thr Leu Leu Thr Leu Lys Asn Leu Thr
    380                 385                 390

Asn Ile Asp Ile Ser Lys Asn Ser Phe His Ser Met Pro Glu Thr Cys
395                 400                 405                 410

Gln Trp Pro Glu Lys Met Lys Tyr Leu Asn Leu Ser Ser Thr Arg Ile
                415                 420                 425

His Ser Val Thr Gly Cys Ile Pro Lys Thr Leu Glu Ile Leu Asp Val
            430                 435                 440
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Ser Asn Asn Asn Leu Asn Leu Phe Ser Leu Asn Leu Pro Gln Leu Lys
445 450 455

Glu Leu Tyr Ile Ser Arg Asn Lys Leu Met Thr Leu Pro Asp Ala Ser
460 465 470

Leu Leu Pro Met Leu Leu Val Leu Lys Ile Ser Arg Asn Ala Ile Thr
475 480 485 490

Thr Phe Ser Lys Glu Gln Leu Asp Ser Phe His Thr Leu Lys Thr Leu
495 500 505

Glu Ala Gly Gly Asn Asn Phe Ile Cys Ser Cys Glu Phe Leu Ser Phe
510 515 520

Thr Gln Glu Gln Gln Ala Leu Ala Lys Val Leu Ile Asp Trp Pro Ala
525 530 535

Asn Tyr Leu Cys Asp Ser Pro Ser His Val Arg Gly Gln Gln Val Gln
540 545 550

Asp Val Arg Leu Ser Val Ser Glu Cys His Arg Thr Ala Leu Val Ser
555 560 565 570

Gly Met Cys Cys Ala Leu Phe Leu Leu Ile Leu Leu Thr Gly Val Leu
575 580 585

Cys His Arg Phe His Gly Leu Trp Tyr Met Lys Met Met Trp Ala Trp
590 595 600

Leu Gln Ala Lys Arg Lys Pro Arg Lys Ala Pro Ser Arg Asn Ile Cys
605 610 615

Tyr Asp Ala Phe Val Ser Tyr Ser Glu Arg Asp Ala Tyr Trp Val Glu
620 625 630

Asn Leu Met Val Gln Glu Leu Glu Asn Phe Asn Pro Pro Phe Lys Leu
635 640 645 650

Cys Leu His Lys Arg Asp Phe Ile Pro Gly Lys Trp Ile Ile Asp Asn
655 660 665

Ile Ile Asp Ser Ile Glu Lys Ser His Lys Thr Val Phe Val Leu Ser
670 675 680

Glu Asn Phe Val Lys Ser Glu Trp Cys Lys Tyr Glu Leu Asp Phe Ser
685 690 695

His Phe Arg Leu Phe Glu Glu Asn Asn Asp Ala Ala Ile Leu Ile Leu
700 705 710

Leu Glu Pro Ile Glu Lys Lys Ala Ile Pro Gln Arg Phe Cys Lys Leu
715 720 725 730

Arg Lys Ile Met Asn Thr Lys Thr Tyr Leu Glu Trp Pro Met Asp Glu
735 740 745

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Ala Gln Arg Glu Gly Phe Trp Val Asn Leu Arg Ala Ala Ile Lys Ser
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<211> 2715
<212> DNA
<213> Unknown

<220>
<223> Description of Unknown Organism:primate; surmised
      Homo sapiens

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<222> (1)..(2712)

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<221> mat_peptide
<222> (64)..(2712)

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Met Arg Gln Thr Leu Pro Cys Ile Tyr Phe Trp Gly Gly Leu Leu Pro
    -20                 -15                 -10

ttt ggg atg ctg tgt gca tcc tcc acc acc aag tgc act gtt agc cat   96
Phe Gly Met Leu Cys Ala Ser Ser Thr Thr Lys Cys Thr Val Ser His
 -5              -1  1                   5                  10

gaa gtt gct gac tgc agc cac ctg aag ttg act cag gta ccc gat gat   144
Glu Val Ala Asp Cys Ser His Leu Lys Leu Thr Gln Val Pro Asp Asp
             15                  20                  25

cta ccc aca aac ata aca gtg ttg aac ctt acc cat aat caa ctc aga   192
Leu Pro Thr Asn Ile Thr Val Leu Asn Leu Thr His Asn Gln Leu Arg
         30                  35                  40

aga tta cca gcc gcc aac ttc aca agg tat agc cag cta act agc ttg   240
Arg Leu Pro Ala Ala Asn Phe Thr Arg Tyr Ser Gln Leu Thr Ser Leu
     45                  50                  55

gat gta gga ttt aac acc atc tca aaa ctg gag cca gaa ttg tgc cag   288
Asp Val Gly Phe Asn Thr Ile Ser Lys Leu Glu Pro Glu Leu Cys Gln
 60                  65                  70                  75

aaa ctt ccc atg tta aaa gtt ttg aac ctc cag cac aat gag cta tct   336
Lys Leu Pro Met Leu Lys Val Leu Asn Leu Gln His Asn Glu Leu Ser
                 80                  85                  90

caa ctt tct gat aaa acc ttt gcc ttc tgc acg aat ttg act gaa ctc   384
Gln Leu Ser Asp Lys Thr Phe Ala Phe Cys Thr Asn Leu Thr Glu Leu
             95                  100                 105

cat ctc atg tcc aac tca atc cag aaa att aaa aat aat ccc ttt gtc   432
His Leu Met Ser Asn Ser Ile Gln Lys Ile Lys Asn Asn Pro Phe Val
         110                 115                 120
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aag cag aag aat tta atc aca tta gat ctg tct cat aat ggc ttg tca   480
Lys Gln Lys Asn Leu Ile Thr Leu Asp Leu Ser His Asn Gly Leu Ser
    125                 130                 135

tct aca aaa tta gga act cag gtt cag ctg gaa aat ctc caa gag ctt   528
Ser Thr Lys Leu Gly Thr Gln Val Gln Leu Glu Asn Leu Gln Glu Leu
140                 145                 150                 155

cta tta tca aac aat aaa att caa gcg cta aaa agt gaa gaa ctg gat   576
Leu Leu Ser Asn Asn Lys Ile Gln Ala Leu Lys Ser Glu Glu Leu Asp
                160                 165                 170

atc ttt gcc aat tca tct tta aaa aaa tta gag ttg tca tcg aat caa   624
Ile Phe Ala Asn Ser Ser Leu Lys Lys Leu Glu Leu Ser Ser Asn Gln
            175                 180                 185

att aaa gag ttt tct cca ggg tgt ttt cac gca att gga aga tta ttt   672
Ile Lys Glu Phe Ser Pro Gly Cys Phe His Ala Ile Gly Arg Leu Phe
        190                 195                 200

ggc ctc ttt ctg aac aat gtc cag ctg ggt ccc agc ctt aca gag aag   720
Gly Leu Phe Leu Asn Asn Val Gln Leu Gly Pro Ser Leu Thr Glu Lys
    205                 210                 215

cta tgt ttg gaa tta gca aac aca agc att cgg aat ctg tct ctg agt   768
Leu Cys Leu Glu Leu Ala Asn Thr Ser Ile Arg Asn Leu Ser Leu Ser
220                 225                 230                 235

aac agc cag ctg tcc acc acc agc aat aca act ttc ttg gga cta aag   816
Asn Ser Gln Leu Ser Thr Thr Ser Asn Thr Thr Phe Leu Gly Leu Lys
                240                 245                 250

tgg aca aat ctc act atg ctc gat ctt tcc tac aac aac tta aat gtg   864
Trp Thr Asn Leu Thr Met Leu Asp Leu Ser Tyr Asn Asn Leu Asn Val
            255                 260                 265

gtt ggt aac gat tcc ttt gct tgg ctt cca caa cta gaa tat ttc ttc   912
Val Gly Asn Asp Ser Phe Ala Trp Leu Pro Gln Leu Glu Tyr Phe Phe
        270                 275                 280

cta gag tat aat aat ata cag cat ttg ttt tct cac tct ttg cac ggg   960
Leu Glu Tyr Asn Asn Ile Gln His Leu Phe Ser His Ser Leu His Gly
    285                 290                 295

ctt ttc aat gtg agg tac ctg aat ttg aaa cgg tct ttt act aaa caa  1008
Leu Phe Asn Val Arg Tyr Leu Asn Leu Lys Arg Ser Phe Thr Lys Gln
300                 305                 310                 315

agt att tcc ctt gcc tca ctc ccc aag att gat gat ttt tct ttt cag  1056
Ser Ile Ser Leu Ala Ser Leu Pro Lys Ile Asp Asp Phe Ser Phe Gln
                320                 325                 330

tgg cta aaa tgt ttg gag cac ctt aac atg gaa gat aat gat att cca  1104
Trp Leu Lys Cys Leu Glu His Leu Asn Met Glu Asp Asn Asp Ile Pro
            335                 340                 345

ggc ata aaa agc aat atg ttc aca gga ttg ata aac ctg aaa tac tta  1152
Gly Ile Lys Ser Asn Met Phe Thr Gly Leu Ile Asn Leu Lys Tyr Leu
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                350                 355                 360

agt cta tcc aac tcc ttt aca agt ttg cga act ttg aca aat gaa aca   1200
Ser Leu Ser Asn Ser Phe Thr Ser Leu Arg Thr Leu Thr Asn Glu Thr
    365                 370                 375

ttt gta tca ctt gct cat tct ccc tta cac ata ctc aac cta acc aag   1248
Phe Val Ser Leu Ala His Ser Pro Leu His Ile Leu Asn Leu Thr Lys
380                 385                 390                 395

aat aaa atc tca aaa ata gag agt gat gct ttc tct tgg ttg ggc cac   1296
Asn Lys Ile Ser Lys Ile Glu Ser Asp Ala Phe Ser Trp Leu Gly His
                400                 405                 410

cta gaa gta ctt gac ctg ggc ctt aat gaa att ggg caa gaa ctc aca   1344
Leu Glu Val Leu Asp Leu Gly Leu Asn Glu Ile Gly Gln Glu Leu Thr
            415                 420                 425

ggc cag gaa tgg aga ggt cta gaa aat att ttc gaa atc tat ctt tcc   1392
Gly Gln Glu Trp Arg Gly Leu Glu Asn Ile Phe Glu Ile Tyr Leu Ser
        430                 435                 440

tac aac aag tac ctg cag ctg act agg aac tcc ttt gcc ttg gtc cca   1440
Tyr Asn Lys Tyr Leu Gln Leu Thr Arg Asn Ser Phe Ala Leu Val Pro
    445                 450                 455

agc ctt caa cga ctg atg ctc cga agg gtg gcc ctt aaa aat gtg gat   1488
Ser Leu Gln Arg Leu Met Leu Arg Arg Val Ala Leu Lys Asn Val Asp
460                 465                 470                 475

agc tct cct tca cca ttc cag cct ctt cgt aac ttg acc att ctg gat   1536
Ser Ser Pro Ser Pro Phe Gln Pro Leu Arg Asn Leu Thr Ile Leu Asp
                480                 485                 490

cta agc aac aac aac ata gcc aac ata aat gat gac atg ttg gag ggt   1584
Leu Ser Asn Asn Asn Ile Ala Asn Ile Asn Asp Asp Met Leu Glu Gly
            495                 500                 505

ctt gag aaa cta gaa att ctc gat ttg cag cat aac aac tta gca cgg   1632
Leu Glu Lys Leu Glu Ile Leu Asp Leu Gln His Asn Asn Leu Ala Arg
        510                 515                 520

ctc tgg aaa cac gca aac cct ggt ggt ccc att tat ttc cta aag ggt   1680
Leu Trp Lys His Ala Asn Pro Gly Gly Pro Ile Tyr Phe Leu Lys Gly
    525                 530                 535

ctg tct cac ctc cac atc ctt aac ttg gag tcc aac ggc ttt gac gag   1728
Leu Ser His Leu His Ile Leu Asn Leu Glu Ser Asn Gly Phe Asp Glu
540                 545                 550                 555

atc cca gtt gag gtc ttc aag gat tta ttt gaa cta aag atc atc gat   1776
Ile Pro Val Glu Val Phe Lys Asp Leu Phe Glu Leu Lys Ile Ile Asp
                560                 565                 570

tta gga ttg aat aat tta aac aca ctt cca gca tct gtc ttt aat aat   1824
Leu Gly Leu Asn Asn Leu Asn Thr Leu Pro Ala Ser Val Phe Asn Asn
            575                 580                 585
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cag gtg tct cta aag tca ttg aac ctt cag aag aat ctc ata aca tcc  1872
Gln Val Ser Leu Lys Ser Leu Asn Leu Gln Lys Asn Leu Ile Thr Ser
        590                 595                 600

gtt gag aag aag gtt ttc ggg cca gct ttc agg aac ctg act gag tta  1920
Val Glu Lys Lys Val Phe Gly Pro Ala Phe Arg Asn Leu Thr Glu Leu
    605                 610                 615

gat atg cgc ttt aat ccc ttt gat tgc acg tgt gaa agt att gcc tgg  1968
Asp Met Arg Phe Asn Pro Phe Asp Cys Thr Cys Glu Ser Ile Ala Trp
620                 625                 630                 635

ttt gtt aat tgg att aac gag acc cat acc aac atc cct gag ctg tca  2016
Phe Val Asn Trp Ile Asn Glu Thr His Thr Asn Ile Pro Glu Leu Ser
                640                 645                 650

agc cac tac ctt tgc aac act cca cct cac tat cat ggg ttc cca gtg  2064
Ser His Tyr Leu Cys Asn Thr Pro Pro His Tyr His Gly Phe Pro Val
            655                 660                 665

aga ctt ttt gat aca tca tct tgc aaa gac agt gcc ccc ttt gaa ctc  2112
Arg Leu Phe Asp Thr Ser Ser Cys Lys Asp Ser Ala Pro Phe Glu Leu
        670                 675                 680

ttt ttc atg atc aat acc agt atc ctg ttg att ttt atc ttt att gta  2160
Phe Phe Met Ile Asn Thr Ser Ile Leu Leu Ile Phe Ile Phe Ile Val
    685                 690                 695

ctt ctc atc cac ttt gag ggc tgg agg ata tct ttt tat tgg aat gtt  2208
Leu Leu Ile His Phe Glu Gly Trp Arg Ile Ser Phe Tyr Trp Asn Val
700                 705                 710                 715

tca gta cat cga gtt ctt ggt ttc aaa gaa ata gac aga cag aca gaa  2256
Ser Val His Arg Val Leu Gly Phe Lys Glu Ile Asp Arg Gln Thr Glu
                720                 725                 730

cag ttt gaa tat gca gca tat ata att cat gcc tat aaa gat aag gat  2304
Gln Phe Glu Tyr Ala Ala Tyr Ile Ile His Ala Tyr Lys Asp Lys Asp
            735                 740                 745

tgg gtc tgg gaa cat ttc tct tca atg gaa aag gaa gac caa tct ctc  2352
Trp Val Trp Glu His Phe Ser Ser Met Glu Lys Glu Asp Gln Ser Leu
        750                 755                 760

aaa ttt tgt ctg gaa gaa agg gac ttt gag gcg ggt gtt ttt gaa cta  2400
Lys Phe Cys Leu Glu Glu Arg Asp Phe Glu Ala Gly Val Phe Glu Leu
    765                 770                 775

gaa gca att gtt aac agc atc aaa aga agc aga aaa att att ttt gtt  2448
Glu Ala Ile Val Asn Ser Ile Lys Arg Ser Arg Lys Ile Ile Phe Val
780                 785                 790                 795

ata aca cac cat cta tta aaa gac cca tta tgc aaa aga ttc aag gta  2496
Ile Thr His His Leu Leu Lys Asp Pro Leu Cys Lys Arg Phe Lys Val
                800                 805                 810

cat cat gca gtt caa caa gct att gaa caa aat ctg gat tcc att ata  2544
His His Ala Val Gln Gln Ala Ile Glu Gln Asn Leu Asp Ser Ile Ile
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                815                 820                 825

ttg gtt ttc ctt gag gag att cca gat tat aaa ctg aac cat gca ctc   2592
Leu Val Phe Leu Glu Glu Ile Pro Asp Tyr Lys Leu Asn His Ala Leu
 830                 835                 840

tgt ttg cga aga gga atg ttt aaa tct cac tgc atc ttg aac tgg cca   2640
Cys Leu Arg Arg Gly Met Phe Lys Ser His Cys Ile Leu Asn Trp Pro
 845                 850                 855

gtt cag aaa gaa cgg ata ggt gcc ttt cgt cat aaa ttg caa gta gca   2688
Val Gln Lys Glu Arg Ile Gly Ala Phe Arg His Lys Leu Gln Val Ala
 860                 865                 870                 875

ctt gga tcc aaa aac tct gta cat taa                              2715
Leu Gly Ser Lys Asn Ser Val His
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Phe Gly Met Leu Cys Ala Ser Ser Thr Thr Lys Cys Thr Val Ser His
 -5                  -1   1                   5                  10

Glu Val Ala Asp Cys Ser His Leu Lys Leu Thr Gln Val Pro Asp Asp
                15                  20                  25

Leu Pro Thr Asn Ile Thr Val Leu Asn Leu Thr His Asn Gln Leu Arg
            30                  35                  40

Arg Leu Pro Ala Ala Asn Phe Thr Arg Tyr Ser Gln Leu Thr Ser Leu
 45                  50                  55

Asp Val Gly Phe Asn Thr Ile Ser Lys Leu Glu Pro Glu Leu Cys Gln
 60                  65                  70                  75

Lys Leu Pro Met Leu Lys Val Leu Asn Leu Gln His Asn Glu Leu Ser
                80                  85                  90

Gln Leu Ser Asp Lys Thr Phe Ala Phe Cys Thr Asn Leu Thr Glu Leu
                95                 100                 105

His Leu Met Ser Asn Ser Ile Gln Lys Ile Lys Asn Asn Pro Phe Val
            110                 115                 120

Lys Gln Lys Asn Leu Ile Thr Leu Asp Leu Ser His Asn Gly Leu Ser
 125                 130                 135

Ser Thr Lys Leu Gly Thr Gln Val Gln Leu Glu Asn Leu Gln Glu Leu
140                 145                 150                 155
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Leu Leu Ser Asn Asn Lys Ile Gln Ala Leu Lys Ser Glu Glu Leu Asp
160 165 170

Ile Phe Ala Asn Ser Ser Leu Lys Lys Leu Glu Leu Ser Ser Asn Gln
175 180 185

Ile Lys Glu Phe Ser Pro Gly Cys Phe His Ala Ile Gly Arg Leu Phe
190 195 200

Gly Leu Phe Leu Asn Asn Val Gln Leu Gly Pro Ser Leu Thr Glu Lys
205 210 215

Leu Cys Leu Glu Leu Ala Asn Thr Ser Ile Arg Asn Leu Ser Leu Ser
220 225 230 235

Asn Ser Gln Leu Ser Thr Thr Ser Asn Thr Thr Phe Leu Gly Leu Lys
240 245 250

Trp Thr Asn Leu Thr Met Leu Asp Leu Ser Tyr Asn Asn Leu Asn Val
255 260 265

Val Gly Asn Asp Ser Phe Ala Trp Leu Pro Gln Leu Glu Tyr Phe Phe
270 275 280

Leu Glu Tyr Asn Asn Ile Gln His Leu Phe Ser His Ser Leu His Gly
285 290 295

Leu Phe Asn Val Arg Tyr Leu Asn Leu Lys Arg Ser Phe Thr Lys Gln
300 305 310 315

Ser Ile Ser Leu Ala Ser Leu Pro Lys Ile Asp Asp Phe Ser Phe Gln
320 325 330

Trp Leu Lys Cys Leu Glu His Leu Asn Met Glu Asp Asn Asp Ile Pro
335 340 345

Gly Ile Lys Ser Asn Met Phe Thr Gly Leu Ile Asn Leu Lys Tyr Leu
350 355 360

Ser Leu Ser Asn Ser Phe Thr Ser Leu Arg Thr Leu Thr Asn Glu Thr
365 370 375

Phe Val Ser Leu Ala His Ser Pro Leu His Ile Leu Asn Leu Thr Lys
380 385 390 395

Asn Lys Ile Ser Lys Ile Glu Ser Asp Ala Phe Ser Trp Leu Gly His
400 405 410

Leu Glu Val Leu Asp Leu Gly Leu Asn Glu Ile Gly Gln Glu Leu Thr
415 420 425

Gly Gln Glu Trp Arg Gly Leu Glu Asn Ile Phe Glu Ile Tyr Leu Ser
430 435 440

Tyr Asn Lys Tyr Leu Gln Leu Thr Arg Asn Ser Phe Ala Leu Val Pro
445 450 455

Ser Leu Gln Arg Leu Met Leu Arg Arg Val Ala Leu Lys Asn Val Asp

460 465 470 475

Ser Ser Pro Ser Pro Phe Gln Pro Leu Arg Asn Leu Thr Ile Leu Asp
480 485 490

Leu Ser Asn Asn Asn Ile Ala Asn Ile Asn Asp Asp Met Leu Glu Gly
495 500 505

Leu Glu Lys Leu Glu Ile Leu Asp Leu Gln His Asn Asn Leu Ala Arg
510 515 520

Leu Trp Lys His Ala Asn Pro Gly Gly Pro Ile Tyr Phe Leu Lys Gly
525 530 535

Leu Ser His Leu His Ile Leu Asn Leu Glu Ser Asn Gly Phe Asp Glu
540 545 550 555

Ile Pro Val Glu Val Phe Lys Asp Leu Phe Glu Leu Lys Ile Ile Asp
560 565 570

Leu Gly Leu Asn Asn Leu Asn Thr Leu Pro Ala Ser Val Phe Asn Asn
575 580 585

Gln Val Ser Leu Lys Ser Leu Asn Leu Gln Lys Asn Leu Ile Thr Ser
590 595 600

Val Glu Lys Lys Val Phe Gly Pro Ala Phe Arg Asn Leu Thr Glu Leu
605 610 615

Asp Met Arg Phe Asn Pro Phe Asp Cys Thr Cys Glu Ser Ile Ala Trp
620 625 630 635

Phe Val Asn Trp Ile Asn Glu Thr His Thr Asn Ile Pro Glu Leu Ser
640 645 650

Ser His Tyr Leu Cys Asn Thr Pro Pro His Tyr His Gly Phe Pro Val
655 660 665

Arg Leu Phe Asp Thr Ser Ser Cys Lys Asp Ser Ala Pro Phe Glu Leu
670 675 680

Phe Phe Met Ile Asn Thr Ser Ile Leu Leu Ile Phe Ile Phe Ile Val
685 690 695

Leu Leu Ile His Phe Glu Gly Trp Arg Ile Ser Phe Tyr Trp Asn Val
700 705 710 715

Ser Val His Arg Val Leu Gly Phe Lys Glu Ile Asp Arg Gln Thr Glu
720 725 730

Gln Phe Glu Tyr Ala Ala Tyr Ile Ile His Ala Tyr Lys Asp Lys Asp
735 740 745

Trp Val Trp Glu His Phe Ser Ser Met Glu Lys Glu Asp Gln Ser Leu
750 755 760

Lys Phe Cys Leu Glu Glu Arg Asp Phe Glu Ala Gly Val Phe Glu Leu
765 770 775

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Glu Ala Ile Val Asn Ser Ile Lys Arg Ser Arg Lys Ile Ile Phe Val
780                 785                 790                 795

Ile Thr His His Leu Leu Lys Asp Pro Leu Cys Lys Arg Phe Lys Val
                800                 805                 810

His His Ala Val Gln Gln Ala Ile Glu Gln Asn Leu Asp Ser Ile Ile
            815                 820                 825

Leu Val Phe Leu Glu Glu Ile Pro Asp Tyr Lys Leu Asn His Ala Leu
        830                 835                 840

Cys Leu Arg Arg Gly Met Phe Lys Ser His Cys Ile Leu Asn Trp Pro
    845                 850                 855

Val Gln Lys Glu Arg Ile Gly Ala Phe Arg His Lys Leu Gln Val Ala
860                 865                 870                 875

Leu Gly Ser Lys Asn Ser Val His
                880
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<213> Unknown

<220>
<223> Description of Unknown Organism:primate; surmised Homo sapiens

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<221> CDS
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<400> 7

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 1               5                  10                  15

aag aac ctg gac ctg agc ttt aat ccc ctg agg cat tta ggc agc tat   96
Lys Asn Leu Asp Leu Ser Phe Asn Pro Leu Arg His Leu Gly Ser Tyr
            20                  25                  30

agc ttc ttc agt ttc cca gaa ctg cag gtg ctg gat tta tcc agg tgt   144
Ser Phe Phe Ser Phe Pro Glu Leu Gln Val Leu Asp Leu Ser Arg Cys
        35                  40                  45

gaa atc cag aca att gaa gat ggg gca tat cag agc cta agc cac ctc   192
Glu Ile Gln Thr Ile Glu Asp Gly Ala Tyr Gln Ser Leu Ser His Leu
    50                  55                  60

tct acc tta ata ttg aca gga aac ccc atc cag agt tta gcc ctg gga   240
Ser Thr Leu Ile Leu Thr Gly Asn Pro Ile Gln Ser Leu Ala Leu Gly
65                  70                  75                  80

gcc ttt tct gga cta tca agt tta cag aag ctg gtg gct gtg gag aca   288
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Ala Phe Ser Gly Leu Ser Ser Leu Gln Lys Leu Val Ala Val Glu Thr
                  85                  90                  95

aat cta gca tct cta gag aac ttc ccc att gga cat ctc aaa act ttg   336
Asn Leu Ala Ser Leu Glu Asn Phe Pro Ile Gly His Leu Lys Thr Leu
            100                 105                 110

aaa gaa ctt aat gtg gct cac aat ctt atc caa tct ttc aaa tta cct   384
Lys Glu Leu Asn Val Ala His Asn Leu Ile Gln Ser Phe Lys Leu Pro
        115                 120                 125

gag tat ttt tct aat ctg acc aat cta gag cac ttg gac ctt tcc agc   432
Glu Tyr Phe Ser Asn Leu Thr Asn Leu Glu His Leu Asp Leu Ser Ser
    130                 135                 140

aac aag att caa agt att tat tgc aca gac ttg cgg gtt cta cat caa   480
Asn Lys Ile Gln Ser Ile Tyr Cys Thr Asp Leu Arg Val Leu His Gln
145                 150                 155                 160

atg ccc cta ctc aat ctc tct tta gac ctg tcc ctg aac cct atg aac   528
Met Pro Leu Leu Asn Leu Ser Leu Asp Leu Ser Leu Asn Pro Met Asn
                165                 170                 175

ttt atc caa cca ggt gca ttt aaa gaa att agg ctt cat aag ctg act   576
Phe Ile Gln Pro Gly Ala Phe Lys Glu Ile Arg Leu His Lys Leu Thr
            180                 185                 190

tta aga aat aat ttt gat agt tta aat gta atg aaa act tgt att caa   624
Leu Arg Asn Asn Phe Asp Ser Leu Asn Val Met Lys Thr Cys Ile Gln
        195                 200                 205

ggt ctg gct ggt tta gaa gtc cat cgt ttg gtt ctg gga gaa ttt aga   672
Gly Leu Ala Gly Leu Glu Val His Arg Leu Val Leu Gly Glu Phe Arg
    210                 215                 220

aat gaa gga aac ttg gaa aag ttt gac aaa tct gct cta gag ggc ctg   720
Asn Glu Gly Asn Leu Glu Lys Phe Asp Lys Ser Ala Leu Glu Gly Leu
225                 230                 235                 240

tgc aat ttg acc att gaa gaa ttc cga tta gca tac tta gac tac tac   768
Cys Asn Leu Thr Ile Glu Glu Phe Arg Leu Ala Tyr Leu Asp Tyr Tyr
                245                 250                 255

ctc gat gat att att gac tta ttt aat tgt ttg aca aat gtt tct tca   816
Leu Asp Asp Ile Ile Asp Leu Phe Asn Cys Leu Thr Asn Val Ser Ser
            260                 265                 270

ttt tcc ctg gtg agt gtg act att gaa agg gta aaa gac ttt tct tat   864
Phe Ser Leu Val Ser Val Thr Ile Glu Arg Val Lys Asp Phe Ser Tyr
        275                 280                 285

aat ttc gga tgg caa cat tta gaa tta gtt aac tgt aaa ttt gga cag   912
Asn Phe Gly Trp Gln His Leu Glu Leu Val Asn Cys Lys Phe Gly Gln
    290                 295                 300

ttt ccc aca ttg aaa ctc aaa tct ctc aaa agg ctt act ttc act tcc   960
Phe Pro Thr Leu Lys Leu Lys Ser Leu Lys Arg Leu Thr Phe Thr Ser
305                 310                 315                 320
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aac aaa ggt ggg aat gct ttt tca gaa gtt gat cta cca agc ctt gag   1008
Asn Lys Gly Gly Asn Ala Phe Ser Glu Val Asp Leu Pro Ser Leu Glu
                325                 330                 335

ttt cta gat ctc agt aga aat ggc ttg agt ttc aaa ggt tgc tgt tct   1056
Phe Leu Asp Leu Ser Arg Asn Gly Leu Ser Phe Lys Gly Cys Cys Ser
            340                 345                 350

caa agt gat ttt ggg aca acc agc cta aag tat tta gat ctg agc ttc   1104
Gln Ser Asp Phe Gly Thr Thr Ser Leu Lys Tyr Leu Asp Leu Ser Phe
        355                 360                 365

aat ggt gtt att acc atg agt tca aac ttc ttg ggc tta gaa caa cta   1152
Asn Gly Val Ile Thr Met Ser Ser Asn Phe Leu Gly Leu Glu Gln Leu
    370                 375                 380

gaa cat ctg gat ttc cag cat tcc aat ttg aaa caa atg agt gag ttt   1200
Glu His Leu Asp Phe Gln His Ser Asn Leu Lys Gln Met Ser Glu Phe
385                 390                 395                 400

tca gta ttc cta tca ctc aga aac ctc att tac ctt gac att tct cat   1248
Ser Val Phe Leu Ser Leu Arg Asn Leu Ile Tyr Leu Asp Ile Ser His
                405                 410                 415

act cac acc aga gtt gct ttc aat ggc atc ttc aat ggc ttg tcc agt   1296
Thr His Thr Arg Val Ala Phe Asn Gly Ile Phe Asn Gly Leu Ser Ser
            420                 425                 430

ctc gaa gtc ttg aaa atg gct ggc aat tct ttc cag gaa aac ttc ctt   1344
Leu Glu Val Leu Lys Met Ala Gly Asn Ser Phe Gln Glu Asn Phe Leu
        435                 440                 445

cca gat atc ttc aca gag ctg aga aac ttg acc ttc ctg gac ctc tct   1392
Pro Asp Ile Phe Thr Glu Leu Arg Asn Leu Thr Phe Leu Asp Leu Ser
    450                 455                 460

cag tgt caa ctg gag cag ttg tct cca aca gca ttt aac tca ctc tcc   1440
Gln Cys Gln Leu Glu Gln Leu Ser Pro Thr Ala Phe Asn Ser Leu Ser
465                 470                 475                 480

agt ctt cag gta cta aat atg agc cac aac aac ttc ttt tca ttg gat   1488
Ser Leu Gln Val Leu Asn Met Ser His Asn Asn Phe Phe Ser Leu Asp
                485                 490                 495

acg ttt cct tat aag tgt ctg aac tcc ctc cag gtt ctt gat tac agt   1536
Thr Phe Pro Tyr Lys Cys Leu Asn Ser Leu Gln Val Leu Asp Tyr Ser
            500                 505                 510

ctc aat cac ata atg act tcc aaa aaa cag gaa cta cag cat ttt cca   1584
Leu Asn His Ile Met Thr Ser Lys Lys Gln Glu Leu Gln His Phe Pro
        515                 520                 525

agt agt cta gct ttc tta aat ctt act cag aat gac ttt gct tgt act   1632
Ser Ser Leu Ala Phe Leu Asn Leu Thr Gln Asn Asp Phe Ala Cys Thr
    530                 535                 540

tgt gaa cac cag agt ttc ctg caa tgg atc aag gac cag agg cag ctc   1680
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Cys Glu His Gln Ser Phe Leu Gln Trp Ile Lys Asp Gln Arg Gln Leu
545                 550                 555                 560

ttg gtg gaa gtt gaa cga atg gaa tgt gca aca cct tca gat aag cag   1728
Leu Val Glu Val Glu Arg Met Glu Cys Ala Thr Pro Ser Asp Lys Gln
                565                 570                 575

ggc atg cct gtg ctg agt ttg aat atc acc tgt cag atg aat aag acc   1776
Gly Met Pro Val Leu Ser Leu Asn Ile Thr Cys Gln Met Asn Lys Thr
            580                 585                 590

atc att ggt gtg tcg gtc ctc agt gtg ctt gta gta tct gtt gta gca   1824
Ile Ile Gly Val Ser Val Leu Ser Val Leu Val Val Ser Val Val Ala
        595                 600                 605

gtt ctg gtc tat aag ttc tat ttt cac ctg atg ctt ctt gct ggc tgc   1872
Val Leu Val Tyr Lys Phe Tyr Phe His Leu Met Leu Leu Ala Gly Cys
    610                 615                 620

ata aag tat ggt aga ggt gaa aac atc tat gat gcc ttt gtt atc tac   1920
Ile Lys Tyr Gly Arg Gly Glu Asn Ile Tyr Asp Ala Phe Val Ile Tyr
625                 630                 635                 640

tca agc cag gat gag gac tgg gta agg aat gag cta gta aag aat tta   1968
Ser Ser Gln Asp Glu Asp Trp Val Arg Asn Glu Leu Val Lys Asn Leu
                645                 650                 655

gaa gaa ggg gtg cct cca ttt cag ctc tgc ctt cac tac aga gac ttt   2016
Glu Glu Gly Val Pro Pro Phe Gln Leu Cys Leu His Tyr Arg Asp Phe
            660                 665                 670

att ccc ggt gtg gcc att gct gcc aac atc atc cat gaa ggt ttc cat   2064
Ile Pro Gly Val Ala Ile Ala Ala Asn Ile Ile His Glu Gly Phe His
        675                 680                 685

aaa agc cga aag gtg att gtt gtg gtg tcc cag cac ttc atc cag agc   2112
Lys Ser Arg Lys Val Ile Val Val Val Ser Gln His Phe Ile Gln Ser
    690                 695                 700

cgc tgg tgt atc ttt gaa tat gag att gct cag acc tgg cag ttt ctg   2160
Arg Trp Cys Ile Phe Glu Tyr Glu Ile Ala Gln Thr Trp Gln Phe Leu
705                 710                 715                 720

agc agt cgt gct ggt atc atc ttc att gtc ctg cag aag gtg gag aag   2208
Ser Ser Arg Ala Gly Ile Ile Phe Ile Val Leu Gln Lys Val Glu Lys
                725                 730                 735

acc ctg ctc agg cag cag gtg gag ctg tac cgc ctt ctc agc agg aac   2256
Thr Leu Leu Arg Gln Gln Val Glu Leu Tyr Arg Leu Leu Ser Arg Asn
            740                 745                 750

act tac ctg gag tgg gag gac agt gtc ctg ggg cgg cac atc ttc tgg   2304
Thr Tyr Leu Glu Trp Glu Asp Ser Val Leu Gly Arg His Ile Phe Trp
        755                 760                 765

aga cga ctc aga aaa gcc ctg ctg gat ggt aaa tca tgg aat cca gaa   2352
Arg Arg Leu Arg Lys Ala Leu Leu Asp Gly Lys Ser Trp Asn Pro Glu
    770                 775                 780
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gga aca gtg ggt aca gga tgc aat tgg cag gaa gca aca tct atc tga 2400
Gly Thr Val Gly Thr Gly Cys Asn Trp Gln Glu Ala Thr Ser Ile
785 790 795

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<211> 799
<212> PRT
<213> Unknown

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Met Glu Leu Asn Phe Tyr Lys Ile Pro Asp Asn Leu Pro Phe Ser Thr
1 5 10 15

Lys Asn Leu Asp Leu Ser Phe Asn Pro Leu Arg His Leu Gly Ser Tyr
20 25 30

Ser Phe Phe Ser Phe Pro Glu Leu Gln Val Leu Asp Leu Ser Arg Cys
35 40 45

Glu Ile Gln Thr Ile Glu Asp Gly Ala Tyr Gln Ser Leu Ser His Leu
50 55 60

Ser Thr Leu Ile Leu Thr Gly Asn Pro Ile Gln Ser Leu Ala Leu Gly
65 70 75 80

Ala Phe Ser Gly Leu Ser Ser Leu Gln Lys Leu Val Ala Val Glu Thr
85 90 95

Asn Leu Ala Ser Leu Glu Asn Phe Pro Ile Gly His Leu Lys Thr Leu
100 105 110

Lys Glu Leu Asn Val Ala His Asn Leu Ile Gln Ser Phe Lys Leu Pro
115 120 125

Glu Tyr Phe Ser Asn Leu Thr Asn Leu Glu His Leu Asp Leu Ser Ser
130 135 140

Asn Lys Ile Gln Ser Ile Tyr Cys Thr Asp Leu Arg Val Leu His Gln
145 150 155 160

Met Pro Leu Leu Asn Leu Ser Leu Asp Leu Ser Leu Asn Pro Met Asn
165 170 175

Phe Ile Gln Pro Gly Ala Phe Lys Glu Ile Arg Leu His Lys Leu Thr
180 185 190

Leu Arg Asn Asn Phe Asp Ser Leu Asn Val Met Lys Thr Cys Ile Gln
195 200 205

Gly Leu Ala Gly Leu Glu Val His Arg Leu Val Leu Gly Glu Phe Arg
210 215 220

Asn Glu Gly Asn Leu Glu Lys Phe Asp Lys Ser Ala Leu Glu Gly Leu
225 230 235 240

Cys Asn Leu Thr Ile Glu Glu Phe Arg Leu Ala Tyr Leu Asp Tyr Tyr

245 250 255

Leu Asp Asp Ile Ile Asp Leu Phe Asn Cys Leu Thr Asn Val Ser Ser
260 265 270

Phe Ser Leu Val Ser Val Thr Ile Glu Arg Val Lys Asp Phe Ser Tyr
275 280 285

Asn Phe Gly Trp Gln His Leu Glu Leu Val Asn Cys Lys Phe Gly Gln
290 295 300

Phe Pro Thr Leu Lys Leu Lys Ser Leu Lys Arg Leu Thr Phe Thr Ser
305 310 315 320

Asn Lys Gly Gly Asn Ala Phe Ser Glu Val Asp Leu Pro Ser Leu Glu
325 330 335

Phe Leu Asp Leu Ser Arg Asn Gly Leu Ser Phe Lys Gly Cys Cys Ser
340 345 350

Gln Ser Asp Phe Gly Thr Thr Ser Leu Lys Tyr Leu Asp Leu Ser Phe
355 360 365

Asn Gly Val Ile Thr Met Ser Ser Asn Phe Leu Gly Leu Glu Gln Leu
370 375 380

Glu His Leu Asp Phe Gln His Ser Asn Leu Lys Gln Met Ser Glu Phe
385 390 395 400

Ser Val Phe Leu Ser Leu Arg Asn Leu Ile Tyr Leu Asp Ile Ser His
405 410 415

Thr His Thr Arg Val Ala Phe Asn Gly Ile Phe Asn Gly Leu Ser Ser
420 425 430

Leu Glu Val Leu Lys Met Ala Gly Asn Ser Phe Gln Glu Asn Phe Leu
435 440 445

Pro Asp Ile Phe Thr Glu Leu Arg Asn Leu Thr Phe Leu Asp Leu Ser
450 455 460

Gln Cys Gln Leu Glu Gln Leu Ser Pro Thr Ala Phe Asn Ser Leu Ser
465 470 475 480

Ser Leu Gln Val Leu Asn Met Ser His Asn Asn Phe Phe Ser Leu Asp
485 490 495

Thr Phe Pro Tyr Lys Cys Leu Asn Ser Leu Gln Val Leu Asp Tyr Ser
500 505 510

Leu Asn His Ile Met Thr Ser Lys Lys Gln Glu Leu Gln His Phe Pro
515 520 525

Ser Ser Leu Ala Phe Leu Asn Leu Thr Gln Asn Asp Phe Ala Cys Thr
530 535 540

Cys Glu His Gln Ser Phe Leu Gln Trp Ile Lys Asp Gln Arg Gln Leu
545 550 555 560

Leu Val Glu Val Glu Arg Met Glu Cys Ala Thr Pro Ser Asp Lys Gln
565 570 575

Gly Met Pro Val Leu Ser Leu Asn Ile Thr Cys Gln Met Asn Lys Thr
580 585 590

Ile Ile Gly Val Ser Val Leu Ser Val Leu Val Val Ser Val Val Ala
595 600 605

Val Leu Val Tyr Lys Phe Tyr Phe His Leu Met Leu Leu Ala Gly Cys
610 615 620

Ile Lys Tyr Gly Arg Gly Glu Asn Ile Tyr Asp Ala Phe Val Ile Tyr
625 630 635 640

Ser Ser Gln Asp Glu Asp Trp Val Arg Asn Glu Leu Val Lys Asn Leu
645 650 655

Glu Glu Gly Val Pro Pro Phe Gln Leu Cys Leu His Tyr Arg Asp Phe
660 665 670

Ile Pro Gly Val Ala Ile Ala Ala Asn Ile Ile His Glu Gly Phe His
675 680 685

Lys Ser Arg Lys Val Ile Val Val Val Ser Gln His Phe Ile Gln Ser
690 695 700

Arg Trp Cys Ile Phe Glu Tyr Glu Ile Ala Gln Thr Trp Gln Phe Leu
705 710 715 720

Ser Ser Arg Ala Gly Ile Ile Phe Ile Val Leu Gln Lys Val Glu Lys
725 730 735

Thr Leu Leu Arg Gln Gln Val Glu Leu Tyr Arg Leu Leu Ser Arg Asn
740 745 750

Thr Tyr Leu Glu Trp Glu Asp Ser Val Leu Gly Arg His Ile Phe Trp
755 760 765

Arg Arg Leu Arg Lys Ala Leu Leu Asp Gly Lys Ser Trp Asn Pro Glu
770 775 780

Gly Thr Val Gly Thr Gly Cys Asn Trp Gln Glu Ala Thr Ser Ile
785 790 795

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<221> CDS

<222> (1)..(1095)

<400> 9

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tgt tgg gat gtt ttt gag gga ctt tct cat ctt caa gtt ctg tat ttg   48
Cys Trp Asp Val Phe Glu Gly Leu Ser His Leu Gln Val Leu Tyr Leu
 1               5                  10                  15

aat cat aac tat ctt aat tcc ctt cca cca gga gta ttt agc cat ctg   96
Asn His Asn Tyr Leu Asn Ser Leu Pro Pro Gly Val Phe Ser His Leu
            20                  25                  30

act gca tta agg gga cta agc ctc aac tcc aac agg ctg aca gtt ctt  144
Thr Ala Leu Arg Gly Leu Ser Leu Asn Ser Asn Arg Leu Thr Val Leu
        35                  40                  45

tct cac aat gat tta cct gct aat tta gag atc ctg gac ata tcc agg  192
Ser His Asn Asp Leu Pro Ala Asn Leu Glu Ile Leu Asp Ile Ser Arg
    50                  55                  60

aac cag ctc cta gct cct aat cct gat gta ttt gta tca ctt agt gtc  240
Asn Gln Leu Leu Ala Pro Asn Pro Asp Val Phe Val Ser Leu Ser Val
65                  70                  75                  80

ttg gat ata act cat aac aag ttc att tgt gaa tgt gaa ctt agc act  288
Leu Asp Ile Thr His Asn Lys Phe Ile Cys Glu Cys Glu Leu Ser Thr
                85                  90                  95

ttt atc aat tgg ctt aat cac acc aat gtc act ata gct ggg cct cct  336
Phe Ile Asn Trp Leu Asn His Thr Asn Val Thr Ile Ala Gly Pro Pro
            100                 105                 110

gca gac ata tat tgt gtg tac cct gac tcg ttc tct ggg gtt tcc ctc  384
Ala Asp Ile Tyr Cys Val Tyr Pro Asp Ser Phe Ser Gly Val Ser Leu
        115                 120                 125

ttc tct ctt tcc acg gaa ggt tgt gat gaa gag gaa gtc tta aag tcc  432
Phe Ser Leu Ser Thr Glu Gly Cys Asp Glu Glu Glu Val Leu Lys Ser
    130                 135                 140

cta aag ttc tcc ctt ttc att gta tgc act gtc act ctg act ctg ttc  480
Leu Lys Phe Ser Leu Phe Ile Val Cys Thr Val Thr Leu Thr Leu Phe
145                 150                 155                 160

ctc atg acc atc ctc aca gtc aca aag ttc cgg ggc ttc tgt ttt atc  528
Leu Met Thr Ile Leu Thr Val Thr Lys Phe Arg Gly Phe Cys Phe Ile
                165                 170                 175

tgt tat aag aca gcc cag aga ctg gtg ttc aag gac cat ccc cag ggc  576
Cys Tyr Lys Thr Ala Gln Arg Leu Val Phe Lys Asp His Pro Gln Gly
            180                 185                 190

aca gaa cct gat atg tac aaa tat gat gcc tat ttg tgc ttc agc agc  624
Thr Glu Pro Asp Met Tyr Lys Tyr Asp Ala Tyr Leu Cys Phe Ser Ser
        195                 200                 205

aaa gac ttc aca tgg gtg cag aat gct ttg ctc aaa cac ctg gac act  672
Lys Asp Phe Thr Trp Val Gln Asn Ala Leu Leu Lys His Leu Asp Thr
    210                 215                 220
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caa tac agt gac caa aac aga ttc aac ctg tgc ttt gaa gaa aga gac 720
Gln Tyr Ser Asp Gln Asn Arg Phe Asn Leu Cys Phe Glu Glu Arg Asp
225 230 235 240

ttt gtc cca gga gaa aac cgc att gcc aat atc cag gat gcc atc tgg 768
Phe Val Pro Gly Glu Asn Arg Ile Ala Asn Ile Gln Asp Ala Ile Trp
245 250 255

aac agt aga aag atc gtt tgt ctt gtg agc aga cac ttc ctt aga gat 816
Asn Ser Arg Lys Ile Val Cys Leu Val Ser Arg His Phe Leu Arg Asp
260 265 270

ggc tgg tgc ctt gaa gcc ttc agt tat gcc cag ggc agg tgc tta tct 864
Gly Trp Cys Leu Glu Ala Phe Ser Tyr Ala Gln Gly Arg Cys Leu Ser
275 280 285

gac ctt aac agt gct ctc atc atg gtg gtg gtt ggg tcc ttg tcc cag 912
Asp Leu Asn Ser Ala Leu Ile Met Val Val Val Gly Ser Leu Ser Gln
290 295 300

tac cag ttg atg aaa cat caa tcc atc aga ggc ttt gta cag aaa cag 960
Tyr Gln Leu Met Lys His Gln Ser Ile Arg Gly Phe Val Gln Lys Gln
305 310 315 320

cag tat ttg agg tgg cct gag gat ctc cag gat gtt ggc tgg ttt ctt 1008
Gln Tyr Leu Arg Trp Pro Glu Asp Leu Gln Asp Val Gly Trp Phe Leu
325 330 335

cat aaa ctc tct caa cag ata cta aag aaa gaa aag gaa aag aag aaa 1056
His Lys Leu Ser Gln Gln Ile Leu Lys Lys Glu Lys Glu Lys Lys Lys
340 345 350

gac aat aac att ccg ttg caa act gta gca acc atc tcc taatcaaagg 1105
Asp Asn Asn Ile Pro Leu Gln Thr Val Ala Thr Ile Ser
355 360 365

agcaatttcc aacttatctc aagccacaaa taactcttca ctttgtattt gcaccaagtt 1165

atcattttgg ggtcctctct ggaggttttt ttttttcttt tgctactatg aaaacaacat 1225

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20 25 30

Thr Ala Leu Arg Gly Leu Ser Leu Asn Ser Asn Arg Leu Thr Val Leu
35 40 45

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Ser His Asn Asp Leu Pro Ala Asn Leu Glu Ile Leu Asp Ile Ser Arg
     50                  55                  60

Asn Gln Leu Leu Ala Pro Asn Pro Asp Val Phe Val Ser Leu Ser Val
 65                  70                  75                  80

Leu Asp Ile Thr His Asn Lys Phe Ile Cys Glu Cys Glu Leu Ser Thr
                 85                  90                  95

Phe Ile Asn Trp Leu Asn His Thr Asn Val Thr Ile Ala Gly Pro Pro
            100                 105                 110

Ala Asp Ile Tyr Cys Val Tyr Pro Asp Ser Phe Ser Gly Val Ser Leu
        115                 120                 125

Phe Ser Leu Ser Thr Glu Gly Cys Asp Glu Glu Glu Val Leu Lys Ser
    130                 135                 140

Leu Lys Phe Ser Leu Phe Ile Val Cys Thr Val Thr Leu Thr Leu Phe
145                 150                 155                 160

Leu Met Thr Ile Leu Thr Val Thr Lys Phe Arg Gly Phe Cys Phe Ile
                165                 170                 175

Cys Tyr Lys Thr Ala Gln Arg Leu Val Phe Lys Asp His Pro Gln Gly
            180                 185                 190

Thr Glu Pro Asp Met Tyr Lys Tyr Asp Ala Tyr Leu Cys Phe Ser Ser
        195                 200                 205

Lys Asp Phe Thr Trp Val Gln Asn Ala Leu Leu Lys His Leu Asp Thr
    210                 215                 220

Gln Tyr Ser Asp Gln Asn Arg Phe Asn Leu Cys Phe Glu Glu Arg Asp
225                 230                 235                 240

Phe Val Pro Gly Glu Asn Arg Ile Ala Asn Ile Gln Asp Ala Ile Trp
                245                 250                 255

Asn Ser Arg Lys Ile Val Cys Leu Val Ser Arg His Phe Leu Arg Asp
            260                 265                 270

Gly Trp Cys Leu Glu Ala Phe Ser Tyr Ala Gln Gly Arg Cys Leu Ser
        275                 280                 285

Asp Leu Asn Ser Ala Leu Ile Met Val Val Val Gly Ser Leu Ser Gln
    290                 295                 300

Tyr Gln Leu Met Lys His Gln Ser Ile Arg Gly Phe Val Gln Lys Gln
305                 310                 315                 320

Gln Tyr Leu Arg Trp Pro Glu Asp Leu Gln Asp Val Gly Trp Phe Leu
                325                 330                 335

His Lys Leu Ser Gln Gln Ile Leu Lys Lys Glu Lys Glu Lys Lys Lys
            340                 345                 350
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Asp Asn Asn Ile Pro Leu Gln Thr Val Ala Thr Ile Ser
        355                 360                 365


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<222> (1)..(3135)

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<222> (67)..(3135)

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Met Trp Thr Leu Lys Arg Leu Ile Leu Ile Leu Phe Asn Ile Ile Leu
        -20                 -15                 -10

att tcc aaa ctc ctt ggg gct aga tgg ttt cct aaa act ctg ccc tgt   96
Ile Ser Lys Leu Leu Gly Ala Arg Trp Phe Pro Lys Thr Leu Pro Cys
     -5              -1   1               5                  10

gat gtc act ctg gat gtt cca aag aac cat gtg atc gtg gac tgc aca   144
Asp Val Thr Leu Asp Val Pro Lys Asn His Val Ile Val Asp Cys Thr
                15                  20                  25

gac aag cat ttg aca gaa att cct gga ggt att ccc acg aac acc acg   192
Asp Lys His Leu Thr Glu Ile Pro Gly Gly Ile Pro Thr Asn Thr Thr
            30                  35                  40

aac ctc acc ctc acc att aac cac ata cca gac atc tcc cca gcg tcc   240
Asn Leu Thr Leu Thr Ile Asn His Ile Pro Asp Ile Ser Pro Ala Ser
        45                  50                  55

ttt cac aga ctg gac cat ctg gta gag atc gat ttc aga tgc aac tgt   288
Phe His Arg Leu Asp His Leu Val Glu Ile Asp Phe Arg Cys Asn Cys
    60                  65                  70

gta cct att cca ctg ggg tca aaa aac aac atg tgc atc aag agg ctg   336
Val Pro Ile Pro Leu Gly Ser Lys Asn Asn Met Cys Ile Lys Arg Leu
 75                  80                  85                  90

cag att aaa ccc aga agc ttt agt gga ctc act tat tta aaa tcc ctt   384
Gln Ile Lys Pro Arg Ser Phe Ser Gly Leu Thr Tyr Leu Lys Ser Leu
                95                 100                 105

tac ctg gat gga aac cag cta cta gag ata ccg cag ggc ctc ccg cct   432
Tyr Leu Asp Gly Asn Gln Leu Leu Glu Ile Pro Gln Gly Leu Pro Pro
            110                 115                 120
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agc tta cag ctt ctc agc ctt gag gcc aac aac atc ttt tcc atc aga   480
Ser Leu Gln Leu Leu Ser Leu Glu Ala Asn Asn Ile Phe Ser Ile Arg
        125                 130                 135

aaa gag aat cta aca gaa ctg gcc aac ata gaa ata ctc tac ctg ggc   528
Lys Glu Asn Leu Thr Glu Leu Ala Asn Ile Glu Ile Leu Tyr Leu Gly
    140                 145                 150

caa aac tgt tat tat cga aat cct tgt tat gtt tca tat tca ata gag   576
Gln Asn Cys Tyr Tyr Arg Asn Pro Cys Tyr Val Ser Tyr Ser Ile Glu
155                 160                 165                 170

aaa gat gcc ttc cta aac ttg aca aag tta aaa gtg ctc tcc ctg aaa   624
Lys Asp Ala Phe Leu Asn Leu Thr Lys Leu Lys Val Leu Ser Leu Lys
                175                 180                 185

gat aac aat gtc aca gcc gtc cct act gtt ttg cca tct act tta aca   672
Asp Asn Asn Val Thr Ala Val Pro Thr Val Leu Pro Ser Thr Leu Thr
            190                 195                 200

gaa cta tat ctc tac aac aac atg att gca aaa atc caa gaa gat gat   720
Glu Leu Tyr Leu Tyr Asn Asn Met Ile Ala Lys Ile Gln Glu Asp Asp
        205                 210                 215

ttt aat aac ctc aac caa tta caa att ctt gac cta agt gga aat tgc   768
Phe Asn Asn Leu Asn Gln Leu Gln Ile Leu Asp Leu Ser Gly Asn Cys
    220                 225                 230

cct cgt tgt tat aat gcc cca ttt cct tgt gcg ccg tgt aaa aat aat   816
Pro Arg Cys Tyr Asn Ala Pro Phe Pro Cys Ala Pro Cys Lys Asn Asn
235                 240                 245                 250

tct ccc cta cag atc cct gta aat gct ttt gat gcg ctg aca gaa tta   864
Ser Pro Leu Gln Ile Pro Val Asn Ala Phe Asp Ala Leu Thr Glu Leu
                255                 260                 265

aaa gtt tta cgt cta cac agt aac tct ctt cag cat gtg ccc cca aga   912
Lys Val Leu Arg Leu His Ser Asn Ser Leu Gln His Val Pro Pro Arg
            270                 275                 280

tgg ttt aag aac atc aac aaa ctc cag gaa ctg gat ctg tcc caa aac   960
Trp Phe Lys Asn Ile Asn Lys Leu Gln Glu Leu Asp Leu Ser Gln Asn
        285                 290                 295

ttc ttg gcc aaa gaa att ggg gat gct aaa ttt ctg cat ttt ctc ccc  1008
Phe Leu Ala Lys Glu Ile Gly Asp Ala Lys Phe Leu His Phe Leu Pro
    300                 305                 310

agc ctc atc caa ttg gat ctg tct ttc aat ttt gaa ctt cag gtc tat  1056
Ser Leu Ile Gln Leu Asp Leu Ser Phe Asn Phe Glu Leu Gln Val Tyr
315                 320                 325                 330

cgt gca tct atg aat cta tca caa gca ttt tct tca ctg aaa agc ctg  1104
Arg Ala Ser Met Asn Leu Ser Gln Ala Phe Ser Ser Leu Lys Ser Leu
                335                 340                 345

aaa att ctg cgg atc aga gga tat gtc ttt aaa gag ttg aaa agc ttt  1152
Lys Ile Leu Arg Ile Arg Gly Tyr Val Phe Lys Glu Leu Lys Ser Phe
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                350                 355                 360

aac ctc tcg cca tta cat aat ctt caa aat ctt gaa gtt ctt gat ctt   1200
Asn Leu Ser Pro Leu His Asn Leu Gln Asn Leu Glu Val Leu Asp Leu
365                 370                 375

ggc act aac ttt ata aaa att gct aac ctc agc atg ttt aaa caa ttt   1248
Gly Thr Asn Phe Ile Lys Ile Ala Asn Leu Ser Met Phe Lys Gln Phe
380                 385                 390

aaa aga ctg aaa gtc ata gat ctt tca gtg aat aaa ata tca cct tca   1296
Lys Arg Leu Lys Val Ile Asp Leu Ser Val Asn Lys Ile Ser Pro Ser
395                 400                 405                 410

gga gat tca agt gaa gtt ggc ttc tgc tca aat gcc aga act tct gta   1344
Gly Asp Ser Ser Glu Val Gly Phe Cys Ser Asn Ala Arg Thr Ser Val
415                 420                 425

gaa agt tat gaa ccc cag gtc ctg gaa caa tta cat tat ttc aga tat   1392
Glu Ser Tyr Glu Pro Gln Val Leu Glu Gln Leu His Tyr Phe Arg Tyr
430                 435                 440

gat aag tat gca agg agt tgc aga ttc aaa aac aaa gag gct tct ttc   1440
Asp Lys Tyr Ala Arg Ser Cys Arg Phe Lys Asn Lys Glu Ala Ser Phe
445                 450                 455

atg tct gtt aat gaa agc tgc tac aag tat ggg cag acc ttg gat cta   1488
Met Ser Val Asn Glu Ser Cys Tyr Lys Tyr Gly Gln Thr Leu Asp Leu
460                 465                 470

agt aaa aat agt ata ttt ttt gtc aag tcc tct gat ttt cag cat ctt   1536
Ser Lys Asn Ser Ile Phe Phe Val Lys Ser Ser Asp Phe Gln His Leu
475                 480                 485                 490

tct ttc ctc aaa tgc ctg aat ctg tca gga aat ctc att agc caa act   1584
Ser Phe Leu Lys Cys Leu Asn Leu Ser Gly Asn Leu Ile Ser Gln Thr
495                 500                 505

ctt aat ggc agt gaa ttc caa cct tta gca gag ctg aga tat ttg gac   1632
Leu Asn Gly Ser Glu Phe Gln Pro Leu Ala Glu Leu Arg Tyr Leu Asp
510                 515                 520

ttc tcc aac aac cgg ctt gat tta ctc cat tca aca gca ttt gaa gag   1680
Phe Ser Asn Asn Arg Leu Asp Leu Leu His Ser Thr Ala Phe Glu Glu
525                 530                 535

ctt cac aaa ctg gaa gtt ctg gat ata agc agt aat agc cat tat ttt   1728
Leu His Lys Leu Glu Val Leu Asp Ile Ser Ser Asn Ser His Tyr Phe
540                 545                 550

caa tca gaa gga att act cat atg cta aac ttt acc aag aac cta aag   1776
Gln Ser Glu Gly Ile Thr His Met Leu Asn Phe Thr Lys Asn Leu Lys
555                 560                 565                 570

gtt ctg cag aaa ctg atg atg aac gac aat gac atc tct tcc tcc acc   1824
Val Leu Gln Lys Leu Met Met Asn Asp Asn Asp Ile Ser Ser Ser Thr
575                 580                 585
```

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agc agg acc atg gag agt gag tct ctt aga act ctg gaa ttc aga gga   1872
Ser Arg Thr Met Glu Ser Glu Ser Leu Arg Thr Leu Glu Phe Arg Gly
            590                 595                 600

aat cac tta gat gtt tta tgg aga gaa ggt gat aac aga tac tta caa   1920
Asn His Leu Asp Val Leu Trp Arg Glu Gly Asp Asn Arg Tyr Leu Gln
        605                 610                 615

tta ttc aag aat ctg cta aaa tta gag gaa tta gac atc tct aaa aat   1968
Leu Phe Lys Asn Leu Leu Lys Leu Glu Glu Leu Asp Ile Ser Lys Asn
    620                 625                 630

tcc cta agt ttc ttg cct tct gga gtt ttt gat ggt atg cct cca aat   2016
Ser Leu Ser Phe Leu Pro Ser Gly Val Phe Asp Gly Met Pro Pro Asn
635                 640                 645                 650

cta aag aat ctc tct ttg gcc aaa aat ggg ctc aaa tct ttc agt tgg   2064
Leu Lys Asn Leu Ser Leu Ala Lys Asn Gly Leu Lys Ser Phe Ser Trp
                655                 660                 665

aag aaa ctc cag tgt cta aag aac ctg gaa act ttg gac ctc agc cac   2112
Lys Lys Leu Gln Cys Leu Lys Asn Leu Glu Thr Leu Asp Leu Ser His
            670                 675                 680

aac caa ctg acc act gtc cct gag aga tta tcc aac tgt tcc aga agc   2160
Asn Gln Leu Thr Thr Val Pro Glu Arg Leu Ser Asn Cys Ser Arg Ser
        685                 690                 695

ctc aag aat ctg att ctt aag aat aat caa atc agg agt ctg acg aag   2208
Leu Lys Asn Leu Ile Leu Lys Asn Asn Gln Ile Arg Ser Leu Thr Lys
    700                 705                 710

tat ttt cta caa gat gcc ttc cag ttg cga tat ctg gat ctc agc tca   2256
Tyr Phe Leu Gln Asp Ala Phe Gln Leu Arg Tyr Leu Asp Leu Ser Ser
715                 720                 725                 730

aat aaa atc cag atg atc caa aag acc agc ttc cca gaa aat gtc ctc   2304
Asn Lys Ile Gln Met Ile Gln Lys Thr Ser Phe Pro Glu Asn Val Leu
                735                 740                 745

aac aat ctg aag atg ttg ctt ttg cat cat aat cgg ttt ctg tgc acc   2352
Asn Asn Leu Lys Met Leu Leu Leu His His Asn Arg Phe Leu Cys Thr
            750                 755                 760

tgt gat gct gtg tgg ttt gtc tgg tgg gtt aac cat acg gag gtg act   2400
Cys Asp Ala Val Trp Phe Val Trp Trp Val Asn His Thr Glu Val Thr
        765                 770                 775

att cct tac ctg gcc aca gat gtg act tgt gtg ggg cca gga gca cac   2448
Ile Pro Tyr Leu Ala Thr Asp Val Thr Cys Val Gly Pro Gly Ala His
    780                 785                 790

aag ggc caa agt gtg atc tcc ctg gat ctg tac acc tgt gag tta gat   2496
Lys Gly Gln Ser Val Ile Ser Leu Asp Leu Tyr Thr Cys Glu Leu Asp
795                 800                 805                 810

ctg act aac ctg att ctg ttc tca ctt tcc ata tct gta tct ctc ttt   2544
Leu Thr Asn Leu Ile Leu Phe Ser Leu Ser Ile Ser Val Ser Leu Phe
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                    815                 820                 825

ctc atg gtg atg atg aca gca agt cac ctc tat ttc tgg gat gtg tgg   2592
Leu Met Val Met Met Thr Ala Ser His Leu Tyr Phe Trp Asp Val Trp
                830                 835                 840

tat att tac cat ttc tgt aag gcc aag ata aag ggg tat cag cgt cta   2640
Tyr Ile Tyr His Phe Cys Lys Ala Lys Ile Lys Gly Tyr Gln Arg Leu
            845                 850                 855

ata tca cca gac tgt tgc tat gat gct ttt att gtg tat gac act aaa   2688
Ile Ser Pro Asp Cys Cys Tyr Asp Ala Phe Ile Val Tyr Asp Thr Lys
        860                 865                 870

gac cca gct gtg acc gag tgg gtt ttg gct gag ctg gtg gcc aaa ctg   2736
Asp Pro Ala Val Thr Glu Trp Val Leu Ala Glu Leu Val Ala Lys Leu
875                 880                 885                 890

gaa gac cca aga gag aaa cat ttt aat tta tgt ctc gag gaa agg gac   2784
Glu Asp Pro Arg Glu Lys His Phe Asn Leu Cys Leu Glu Glu Arg Asp
                895                 900                 905

tgg tta cca ggg cag cca gtt ctg gaa aac ctt tcc cag agc ata cag   2832
Trp Leu Pro Gly Gln Pro Val Leu Glu Asn Leu Ser Gln Ser Ile Gln
            910                 915                 920

ctt agc aaa aag aca gtg ttt gtg atg aca gac aag tat gca aag act   2880
Leu Ser Lys Lys Thr Val Phe Val Met Thr Asp Lys Tyr Ala Lys Thr
        925                 930                 935

gaa aat ttt aag ata gca ttt tac ttg tcc cat cag agg ctc atg gat   2928
Glu Asn Phe Lys Ile Ala Phe Tyr Leu Ser His Gln Arg Leu Met Asp
    940                 945                 950

gaa aaa gtt gat gtg att atc ttg ata ttt ctt gag aag ccc ttt cag   2976
Glu Lys Val Asp Val Ile Ile Leu Ile Phe Leu Glu Lys Pro Phe Gln
955                 960                 965                 970

aag tcc aag ttc ctc cag ctc cgg aaa agg ctc tgt ggg agt tct gtc   3024
Lys Ser Lys Phe Leu Gln Leu Arg Lys Arg Leu Cys Gly Ser Ser Val
                975                 980                 985

ctt gag tgg cca aca aac ccg caa gct cac cca tac ttc tgg cag tgt   3072
Leu Glu Trp Pro Thr Asn Pro Gln Ala His Pro Tyr Phe Trp Gln Cys
            990                 995                1000

cta aag aac gcc ctg gcc aca gac aat cat gtg gcc tat agt cag gtg   3120
Leu Lys Asn Ala Leu Ala Thr Asp Asn His Val Ala Tyr Ser Gln Val
        1005                1010                1015

ttc aag gaa acg gtc tag                                          3138
Phe Lys Glu Thr Val
    1020
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<210> 12
<211> 1045
<212> PRT

<213> Unknown

<400> 12

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Met Trp Thr Leu Lys Arg Leu Ile Leu Ile Leu Phe Asn Ile Ile Leu
        -20                 -15                 -10

Ile Ser Lys Leu Leu Gly Ala Arg Trp Phe Pro Lys Thr Leu Pro Cys
     -5              -1  1               5                  10

Asp Val Thr Leu Asp Val Pro Lys Asn His Val Ile Val Asp Cys Thr
                 15                  20                  25

Asp Lys His Leu Thr Glu Ile Pro Gly Gly Ile Pro Thr Asn Thr Thr
             30                  35                  40

Asn Leu Thr Leu Thr Ile Asn His Ile Pro Asp Ile Ser Pro Ala Ser
         45                  50                  55

Phe His Arg Leu Asp His Leu Val Glu Ile Asp Phe Arg Cys Asn Cys
     60                  65                  70

Val Pro Ile Pro Leu Gly Ser Lys Asn Asn Met Cys Ile Lys Arg Leu
 75                  80                  85                  90

Gln Ile Lys Pro Arg Ser Phe Ser Gly Leu Thr Tyr Leu Lys Ser Leu
                 95                 100                 105

Tyr Leu Asp Gly Asn Gln Leu Leu Glu Ile Pro Gln Gly Leu Pro Pro
            110                 115                 120

Ser Leu Gln Leu Leu Ser Leu Glu Ala Asn Asn Ile Phe Ser Ile Arg
        125                 130                 135

Lys Glu Asn Leu Thr Glu Leu Ala Asn Ile Glu Ile Leu Tyr Leu Gly
    140                 145                 150

Gln Asn Cys Tyr Tyr Arg Asn Pro Cys Tyr Val Ser Tyr Ser Ile Glu
155                 160                 165                 170

Lys Asp Ala Phe Leu Asn Leu Thr Lys Leu Lys Val Leu Ser Leu Lys
                175                 180                 185

Asp Asn Asn Val Thr Ala Val Pro Thr Val Leu Pro Ser Thr Leu Thr
            190                 195                 200

Glu Leu Tyr Leu Tyr Asn Asn Met Ile Ala Lys Ile Gln Glu Asp Asp
        205                 210                 215

Phe Asn Asn Leu Asn Gln Leu Gln Ile Leu Asp Leu Ser Gly Asn Cys
    220                 225                 230

Pro Arg Cys Tyr Asn Ala Pro Phe Pro Cys Ala Pro Cys Lys Asn Asn
235                 240                 245                 250

Ser Pro Leu Gln Ile Pro Val Asn Ala Phe Asp Ala Leu Thr Glu Leu
                255                 260                 265

Lys Val Leu Arg Leu His Ser Asn Ser Leu Gln His Val Pro Pro Arg
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270 275 280

Trp Phe Lys Asn Ile Asn Lys Leu Gln Glu Leu Asp Leu Ser Gln Asn
285 290 295

Phe Leu Ala Lys Glu Ile Gly Asp Ala Lys Phe Leu His Phe Leu Pro
300 305 310

Ser Leu Ile Gln Leu Asp Leu Ser Phe Asn Phe Glu Leu Gln Val Tyr
315 320 325 330

Arg Ala Ser Met Asn Leu Ser Gln Ala Phe Ser Ser Leu Lys Ser Leu
335 340 345

Lys Ile Leu Arg Ile Arg Gly Tyr Val Phe Lys Glu Leu Lys Ser Phe
350 355 360

Asn Leu Ser Pro Leu His Asn Leu Gln Asn Leu Glu Val Leu Asp Leu
365 370 375

Gly Thr Asn Phe Ile Lys Ile Ala Asn Leu Ser Met Phe Lys Gln Phe
380 385 390

Lys Arg Leu Lys Val Ile Asp Leu Ser Val Asn Lys Ile Ser Pro Ser
395 400 405 410

Gly Asp Ser Ser Glu Val Gly Phe Cys Ser Asn Ala Arg Thr Ser Val
415 420 425

Glu Ser Tyr Glu Pro Gln Val Leu Glu Gln Leu His Tyr Phe Arg Tyr
430 435 440

Asp Lys Tyr Ala Arg Ser Cys Arg Phe Lys Asn Lys Glu Ala Ser Phe
445 450 455

Met Ser Val Asn Glu Ser Cys Tyr Lys Tyr Gly Gln Thr Leu Asp Leu
460 465 470

Ser Lys Asn Ser Ile Phe Phe Val Lys Ser Ser Asp Phe Gln His Leu
475 480 485 490

Ser Phe Leu Lys Cys Leu Asn Leu Ser Gly Asn Leu Ile Ser Gln Thr
495 500 505

Leu Asn Gly Ser Glu Phe Gln Pro Leu Ala Glu Leu Arg Tyr Leu Asp
510 515 520

Phe Ser Asn Asn Arg Leu Asp Leu Leu His Ser Thr Ala Phe Glu Glu
525 530 535

Leu His Lys Leu Glu Val Leu Asp Ile Ser Ser Asn Ser His Tyr Phe
540 545 550

Gln Ser Glu Gly Ile Thr His Met Leu Asn Phe Thr Lys Asn Leu Lys
555 560 565 570

Val Leu Gln Lys Leu Met Met Asn Asp Asn Asp Ile Ser Ser Ser Thr
575 580 585

Ser Arg Thr Met Glu Ser Glu Ser Leu Arg Thr Leu Glu Phe Arg Gly
590 595 600

Asn His Leu Asp Val Leu Trp Arg Glu Gly Asp Asn Arg Tyr Leu Gln
605 610 615

Leu Phe Lys Asn Leu Leu Lys Leu Glu Glu Leu Asp Ile Ser Lys Asn
620 625 630

Ser Leu Ser Phe Leu Pro Ser Gly Val Phe Asp Gly Met Pro Pro Asn
635 640 645 650

Leu Lys Asn Leu Ser Leu Ala Lys Asn Gly Leu Lys Ser Phe Ser Trp
655 660 665

Lys Lys Leu Gln Cys Leu Lys Asn Leu Glu Thr Leu Asp Leu Ser His
670 675 680

Asn Gln Leu Thr Thr Val Pro Glu Arg Leu Ser Asn Cys Ser Arg Ser
685 690 695

Leu Lys Asn Leu Ile Leu Lys Asn Asn Gln Ile Arg Ser Leu Thr Lys
700 705 710

Tyr Phe Leu Gln Asp Ala Phe Gln Leu Arg Tyr Leu Asp Leu Ser Ser
715 720 725 730

Asn Lys Ile Gln Met Ile Gln Lys Thr Ser Phe Pro Glu Asn Val Leu
735 740 745

Asn Asn Leu Lys Met Leu Leu Leu His His Asn Arg Phe Leu Cys Thr
750 755 760

Cys Asp Ala Val Trp Phe Val Trp Trp Val Asn His Thr Glu Val Thr
765 770 775

Ile Pro Tyr Leu Ala Thr Asp Val Thr Cys Val Gly Pro Gly Ala His
780 785 790

Lys Gly Gln Ser Val Ile Ser Leu Asp Leu Tyr Thr Cys Glu Leu Asp
795 800 805 810

Leu Thr Asn Leu Ile Leu Phe Ser Leu Ser Ile Ser Val Ser Leu Phe
815 820 825

Leu Met Val Met Met Thr Ala Ser His Leu Tyr Phe Trp Asp Val Trp
830 835 840

Tyr Ile Tyr His Phe Cys Lys Ala Lys Ile Lys Gly Tyr Gln Arg Leu
845 850 855

Ile Ser Pro Asp Cys Cys Tyr Asp Ala Phe Ile Val Tyr Asp Thr Lys
860 865 870

Asp Pro Ala Val Thr Glu Trp Val Leu Ala Glu Leu Val Ala Lys Leu
875 880 885 890

Glu Asp Pro Arg Glu Lys His Phe Asn Leu Cys Leu Glu Glu Arg Asp
895 900 905

Trp Leu Pro Gly Gln Pro Val Leu Glu Asn Leu Ser Gln Ser Ile Gln
910 915 920

Leu Ser Lys Lys Thr Val Phe Val Met Thr Asp Lys Tyr Ala Lys Thr
925 930 935

Glu Asn Phe Lys Ile Ala Phe Tyr Leu Ser His Gln Arg Leu Met Asp
940 945 950

Glu Lys Val Asp Val Ile Ile Leu Ile Phe Leu Glu Lys Pro Phe Gln
955 960 965 970

Lys Ser Lys Phe Leu Gln Leu Arg Lys Arg Leu Cys Gly Ser Ser Val
975 980 985

Leu Glu Trp Pro Thr Asn Pro Gln Ala His Pro Tyr Phe Trp Gln Cys
990 995 1000

Leu Lys Asn Ala Leu Ala Thr Asp Asn His Val Ala Tyr Ser Gln Val
1005 1010 1015

Phe Lys Glu Thr Val
1020

<210> 13
<211> 180
<212> DNA
<213> Unknown

<220>
<223> Description of Unknown Organism:rodent; surmised
Mus musculus

<220>
<221> CDS
<222> (1)..(177)

<400> 13
ctt gga aaa cct ctt cag aag tct aag ttt ctt cag ctc agg aag aga 48
Leu Gly Lys Pro Leu Gln Lys Ser Lys Phe Leu Gln Leu Arg Lys Arg
1 5 10 15

ctc tgc agg agc tct gtc ctt gag tgg cct gca aat cca cag gct cac 96
Leu Cys Arg Ser Ser Val Leu Glu Trp Pro Ala Asn Pro Gln Ala His
20 25 30

cca tac ttc tgg cag tgc ctg aaa aat gcc ctg acc aca gac aat cat 144
Pro Tyr Phe Trp Gln Cys Leu Lys Asn Ala Leu Thr Thr Asp Asn His
35 40 45

gtg gct tat agt caa atg ttc aag gaa aca gtc tag 180
Val Ala Tyr Ser Gln Met Phe Lys Glu Thr Val
50 55

<210> 14
<211> 59
<212> PRT
<213> Unknown

<400> 14
Leu Gly Lys Pro Leu Gln Lys Ser Lys Phe Leu Gln Leu Arg Lys Arg
1 5 10 15

Leu Cys Arg Ser Ser Val Leu Glu Trp Pro Ala Asn Pro Gln Ala His
20 25 30

Pro Tyr Phe Trp Gln Cys Leu Lys Asn Ala Leu Thr Thr Asp Asn His
35 40 45

Val Ala Tyr Ser Gln Met Phe Lys Glu Thr Val
50 55

<210> 15
<211> 990
<212> DNA
<213> Unknown

<220>
<223> Description of Unknown Organism:primate; surmised
Homo sapiens

<220>
<221> CDS
<222> (2)..(988)

<400> 15
g aat tcc aga ctt ata aac ttg aaa aat ctc tat ttg gcc tgg aac tgc 49
Asn Ser Arg Leu Ile Asn Leu Lys Asn Leu Tyr Leu Ala Trp Asn Cys
1 5 10 15

tat ttt aac aaa gtt tgc gag aaa act aac ata gaa gat gga gta ttt 97
Tyr Phe Asn Lys Val Cys Glu Lys Thr Asn Ile Glu Asp Gly Val Phe
20 25 30

gaa acg ctg aca aat ttg gag ttg cta tca cta tct ttc aat tct ctt 145
Glu Thr Leu Thr Asn Leu Glu Leu Leu Ser Leu Ser Phe Asn Ser Leu
35 40 45

tca cat gtg cca ccc aaa ctg cca agc tcc cta cgc aaa ctt ttt ctg 193
Ser His Val Pro Pro Lys Leu Pro Ser Ser Leu Arg Lys Leu Phe Leu
50 55 60

agc aac acc cag atc aaa tac att agt gaa gaa gat ttc aag gga ttg 241
Ser Asn Thr Gln Ile Lys Tyr Ile Ser Glu Glu Asp Phe Lys Gly Leu
65 70 75 80

ata aat tta aca tta cta gat tta agc ggg aac tgt ccg agg tgc ttc 289
Ile Asn Leu Thr Leu Leu Asp Leu Ser Gly Asn Cys Pro Arg Cys Phe
85 90 95

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aat gcc cca ttt cca tgc gtg cct tgt gat ggt ggt gct tca att aat   337
Asn Ala Pro Phe Pro Cys Val Pro Cys Asp Gly Gly Ala Ser Ile Asn
            100                 105                 110

ata gat cgt ttt gct ttt caa aac ttg acc caa ctt cga tac cta aac   385
Ile Asp Arg Phe Ala Phe Gln Asn Leu Thr Gln Leu Arg Tyr Leu Asn
        115                 120                 125

ctc tct agc act tcc ctc agg aag att aat gct gcc tgg ttt aaa aat   433
Leu Ser Ser Thr Ser Leu Arg Lys Ile Asn Ala Ala Trp Phe Lys Asn
    130                 135                 140

atg cct cat ctg aag gtg ctg gat ctt gaa ttc aac tat tta gtg gga   481
Met Pro His Leu Lys Val Leu Asp Leu Glu Phe Asn Tyr Leu Val Gly
145                 150                 155                 160

gaa ata gcc tct ggg gca ttt tta acg atg ctg ccc cgc tta gaa ata   529
Glu Ile Ala Ser Gly Ala Phe Leu Thr Met Leu Pro Arg Leu Glu Ile
                165                 170                 175

ctt gac ttg tct ttt aac tat ata aag ggg agt tat cca cag cat att   577
Leu Asp Leu Ser Phe Asn Tyr Ile Lys Gly Ser Tyr Pro Gln His Ile
            180                 185                 190

aat att tcc aga aac ttc tct aaa ctt ttg tct cta cgg gca ttg cat   625
Asn Ile Ser Arg Asn Phe Ser Lys Leu Leu Ser Leu Arg Ala Leu His
        195                 200                 205

tta aga ggt tat gtg ttc cag gaa ctc aga gaa gat gat ttc cag ccc   673
Leu Arg Gly Tyr Val Phe Gln Glu Leu Arg Glu Asp Asp Phe Gln Pro
    210                 215                 220

ctg atg cag ctt cca aac tta tcg act atc aac ttg ggt att aat ttt   721
Leu Met Gln Leu Pro Asn Leu Ser Thr Ile Asn Leu Gly Ile Asn Phe
225                 230                 235                 240

att aag caa atc gat ttc aaa ctt ttc caa aat ttc tcc aat ctg gaa   769
Ile Lys Gln Ile Asp Phe Lys Leu Phe Gln Asn Phe Ser Asn Leu Glu
                245                 250                 255

att att tac ttg tca gaa aac aga ata tca ccg ttg gta aaa gat acc   817
Ile Ile Tyr Leu Ser Glu Asn Arg Ile Ser Pro Leu Val Lys Asp Thr
            260                 265                 270

cgg cag agt tat gca aat agt tcc tct ttt caa cgt cat atc cgg aaa   865
Arg Gln Ser Tyr Ala Asn Ser Ser Ser Phe Gln Arg His Ile Arg Lys
        275                 280                 285

cga cgc tca aca gat ttt gag ttt gac cca cat tcg aac ttt tat cat   913
Arg Arg Ser Thr Asp Phe Glu Phe Asp Pro His Ser Asn Phe Tyr His
    290                 295                 300

ttc acc cgt cct tta ata aag cca caa tgt gct gct tat gga aaa gcc   961
Phe Thr Arg Pro Leu Ile Lys Pro Gln Cys Ala Ala Tyr Gly Lys Ala
305                 310                 315                 320

tta gat tta agc ctc aac agt att ttc tt                           990
Leu Asp Leu Ser Leu Asn Ser Ile Phe
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325

<210> 16
<211> 329
<212> PRT
<213> Unknown

<400> 16

Asn Ser Arg Leu Ile Asn Leu Lys Asn Leu Tyr Leu Ala Trp Asn Cys
1 5 10 15

Tyr Phe Asn Lys Val Cys Glu Lys Thr Asn Ile Glu Asp Gly Val Phe
20 25 30

Glu Thr Leu Thr Asn Leu Glu Leu Leu Ser Leu Ser Phe Asn Ser Leu
35 40 45

Ser His Val Pro Pro Lys Leu Pro Ser Ser Leu Arg Lys Leu Phe Leu
50 55 60

Ser Asn Thr Gln Ile Lys Tyr Ile Ser Glu Glu Asp Phe Lys Gly Leu
65 70 75 80

Ile Asn Leu Thr Leu Leu Asp Leu Ser Gly Asn Cys Pro Arg Cys Phe
85 90 95

Asn Ala Pro Phe Pro Cys Val Pro Cys Asp Gly Gly Ala Ser Ile Asn
100 105 110

Ile Asp Arg Phe Ala Phe Gln Asn Leu Thr Gln Leu Arg Tyr Leu Asn
115 120 125

Leu Ser Ser Thr Ser Leu Arg Lys Ile Asn Ala Ala Trp Phe Lys Asn
130 135 140

Met Pro His Leu Lys Val Leu Asp Leu Glu Phe Asn Tyr Leu Val Gly
145 150 155 160

Glu Ile Ala Ser Gly Ala Phe Leu Thr Met Leu Pro Arg Leu Glu Ile
165 170 175

Leu Asp Leu Ser Phe Asn Tyr Ile Lys Gly Ser Tyr Pro Gln His Ile
180 185 190

Asn Ile Ser Arg Asn Phe Ser Lys Leu Leu Ser Leu Arg Ala Leu His
195 200 205

Leu Arg Gly Tyr Val Phe Gln Glu Leu Arg Glu Asp Asp Phe Gln Pro
210 215 220

Leu Met Gln Leu Pro Asn Leu Ser Thr Ile Asn Leu Gly Ile Asn Phe
225 230 235 240

Ile Lys Gln Ile Asp Phe Lys Leu Phe Gln Asn Phe Ser Asn Leu Glu
245 250 255

Ile Ile Tyr Leu Ser Glu Asn Arg Ile Ser Pro Leu Val Lys Asp Thr

```
                260                 265                 270

Arg Gln Ser Tyr Ala Asn Ser Ser Ser Phe Gln Arg His Ile Arg Lys
        275                 280                 285

Arg Arg Ser Thr Asp Phe Glu Phe Asp Pro His Ser Asn Phe Tyr His
    290                 295                 300

Phe Thr Arg Pro Leu Ile Lys Pro Gln Cys Ala Ala Tyr Gly Lys Ala
305                 310                 315                 320

Leu Asp Leu Ser Leu Asn Ser Ile Phe
                325
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<210> 17
<211> 1557
<212> DNA
<213> Unknown

<220>
<223> Description of Unknown Organism:primate; surmised Homo sapiens

<220>
<221> CDS
<222> (1)..(513)

<220>
<221> misc_feature
<222> (93)..(149)
<223> Xaa translation depends on genetic code

<400> 17

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cag tct ctt tcc aca tcc caa act ttc tat gat gct tac att tct tat   48
Gln Ser Leu Ser Thr Ser Gln Thr Phe Tyr Asp Ala Tyr Ile Ser Tyr
  1               5                  10                  15

gac acc aaa gat gcc tct gtt act gac tgg gtg ata aat gag ctg cgc   96
Asp Thr Lys Asp Ala Ser Val Thr Asp Trp Val Ile Asn Glu Leu Arg
             20                  25                  30

tac cac ctt gaa gag agc cga gac aaa aac gtt ctc ctt tgt cta gag   144
Tyr His Leu Glu Glu Ser Arg Asp Lys Asn Val Leu Leu Cys Leu Glu
         35                  40                  45

gag agg gat tgg gac ccg gga ttg gcc atc atc gac aac ctc atg cag   192
Glu Arg Asp Trp Asp Pro Gly Leu Ala Ile Ile Asp Asn Leu Met Gln
     50                  55                  60

agc atc aac caa agc aag aaa aca gta ttt gtt tta acc aaa aaa tat   240
Ser Ile Asn Gln Ser Lys Lys Thr Val Phe Val Leu Thr Lys Lys Tyr
 65                  70                  75                  80

gca aaa agc tgg aac ttt aaa aca gct ttt tac ttg gsc ttg cag agg   288
Ala Lys Ser Trp Asn Phe Lys Thr Ala Phe Tyr Leu Xaa Leu Gln Arg
                 85                  90                  95
```

cta atg ggt gag aac atg gat gtg att ata ttt atc ctg ctg gag cca 336
Leu Met Gly Glu Asn Met Asp Val Ile Ile Phe Ile Leu Leu Glu Pro
100 105 110

gtg tta cag cat tct ccg tat ttg agg cta cgg cag cgg atc tgt aag 384
Val Leu Gln His Ser Pro Tyr Leu Arg Leu Arg Gln Arg Ile Cys Lys
115 120 125

agc tcc atc ctc cag tgg cct gac aac ccg aag gca gaa agg ttg ttt 432
Ser Ser Ile Leu Gln Trp Pro Asp Asn Pro Lys Ala Glu Arg Leu Phe
130 135 140

tgg caa act ctg wga aat gtg gtc ttg act gaa aat gat tca cgg tat 480
Trp Gln Thr Leu Xaa Asn Val Val Leu Thr Glu Asn Asp Ser Arg Tyr
145 150 155 160

aac aat atg tat gtc gat tcc att aag caa tac taactgacgt taagtcatga 533
Asn Asn Met Tyr Val Asp Ser Ile Lys Gln Tyr
165 170

tttcgcgcca taataaagat gcaaaggaat gacatttcng tattagttat ctattgctan 593

ggtaacnaaa ttantcccaa aaancttang tnggtttnaa aacaacnaca ttntgctggn 653

cccacagttt ttgagggtca ggagtccagg cccagcataa ctgggtcttc tgcttcaggg 713

tgtctncaga ggctgcaatg taggtgttca ccagagacat aggcatcact ggggtcacac 773

tncatgtggt tgttttctgg attcaattcc tcctgggcta ttggccaaag gctatactca 833

tgtaagccat gcgagcctat cccacaangg cagcttgctt catcagagct agcaaaaaag 893

agaggttgct agcaagatga agtcacaatc ttttgtaatc gaatcaaaaa agtgatatct 953

catcactttg gccatattct atttgttaga agtaaaccac aggtcccacc agctccatgg 1013

gagtgaccac ctcagtccag ggaaaacagc tgaagaccaa gatggtgagc tctgattgct 1073

tcagttggtc atcaactatt ttcccttgac tgctgtcctg ggatggccgg ctatcttgat 1133

ggatagattg tgaatatcag gaggccaggg atcactgtgg accatcttag cagttgacct 1193

aacacatctt cttttcaata tctaagaact tttgccactg tgactaatgg tcctaatatt 1253

aagctgttgt ttatatttat catatatcta tggctacatg gttatattat gctgtggttg 1313

cgttcggttt tatttacagt tgcttttaca aatatttgct gtaacatttg acttctaagg 1373

tttagatgcc atttaagaac tgagatggat agcttttaaa gcatctttta cttcttacca 1433

ttttttaaaa gtatgcagct aaattcgaag cttttggtct atattgttaa ttgccattgc 1493

tgtaaatctt aaaatgaatg aataaaaatg tttcatttta aaaaaaaaaa aaaaaaaaaa 1553

aaaa 1557

<210> 18

<211> 171
<212> PRT
<213> Unknown

<400> 18
Gln Ser Leu Ser Thr Ser Gln Thr Phe Tyr Asp Ala Tyr Ile Ser Tyr
1 5 10 15

Asp Thr Lys Asp Ala Ser Val Thr Asp Trp Val Ile Asn Glu Leu Arg
20 25 30

Tyr His Leu Glu Glu Ser Arg Asp Lys Asn Val Leu Leu Cys Leu Glu
35 40 45

Glu Arg Asp Trp Asp Pro Gly Leu Ala Ile Ile Asp Asn Leu Met Gln
50 55 60

Ser Ile Asn Gln Ser Lys Lys Thr Val Phe Val Leu Thr Lys Lys Tyr
65 70 75 80

Ala Lys Ser Trp Asn Phe Lys Thr Ala Phe Tyr Leu Xaa Leu Gln Arg
85 90 95

Leu Met Gly Glu Asn Met Asp Val Ile Ile Phe Ile Leu Leu Glu Pro
100 105 110

Val Leu Gln His Ser Pro Tyr Leu Arg Leu Arg Gln Arg Ile Cys Lys
115 120 125

Ser Ser Ile Leu Gln Trp Pro Asp Asn Pro Lys Ala Glu Arg Leu Phe
130 135 140

Trp Gln Thr Leu Xaa Asn Val Val Leu Thr Glu Asn Asp Ser Arg Tyr
145 150 155 160

Asn Asn Met Tyr Val Asp Ser Ile Lys Gln Tyr
165 170

<210> 19
<211> 629
<212> DNA
<213> Unknown

<220>
<223> Description of Unknown Organism:primate; surmised Homo sapiens

<220>
<221> CDS
<222> (1)..(486)

<220>
<221> misc_feature
<222> (48)..(75)
<223> Xaa translation depends on genetic code

<400> 19

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aat gaa ttg atc ccc aat cta gag aag gaa gat ggt tct atc ttg att   48
Asn Glu Leu Ile Pro Asn Leu Glu Lys Glu Asp Gly Ser Ile Leu Ile
  1               5                  10                  15

tgc ctt tat gaa agc tac ttt gac cct ggc aaa agc att agt gaa aat   96
Cys Leu Tyr Glu Ser Tyr Phe Asp Pro Gly Lys Ser Ile Ser Glu Asn
             20                  25                  30

att gta agc ttc att gag aaa agc tat aag tcc atc ttt gtt ttg tcy   144
Ile Val Ser Phe Ile Glu Lys Ser Tyr Lys Ser Ile Phe Val Leu Xaa
         35                  40                  45

ccc aac ttt gtc cag aat gag tgg tgc cat tat gaa ttc tac ttt gcc   192
Pro Asn Phe Val Gln Asn Glu Trp Cys His Tyr Glu Phe Tyr Phe Ala
     50                  55                  60

cac cac aat ctc ttc cat gaa aat tct gat cay ata att ctt atc tta   240
His His Asn Leu Phe His Glu Asn Ser Asp Xaa Ile Ile Leu Ile Leu
 65                  70                  75                  80

ctg gaa ccc att cca ttc tat tgc att ccc acc agg tat cat aaa ctg   288
Leu Glu Pro Ile Pro Phe Tyr Cys Ile Pro Thr Arg Tyr His Lys Leu
                 85                  90                  95

gaa gct ctc ctg gaa aaa aaa gca tac ttg gaa tgg ccc aag gat agg   336
Glu Ala Leu Leu Glu Lys Lys Ala Tyr Leu Glu Trp Pro Lys Asp Arg
            100                 105                 110

cgt aaa tgt ggg ctt ttc tgg gca aac ctt cga gct gct gtt aat gtt   384
Arg Lys Cys Gly Leu Phe Trp Ala Asn Leu Arg Ala Ala Val Asn Val
        115                 120                 125

aat gta tta gcc acc aga gaa atg tat gaa ctg cag aca ttc aca gag   432
Asn Val Leu Ala Thr Arg Glu Met Tyr Glu Leu Gln Thr Phe Thr Glu
    130                 135                 140

tta aat gaa gag tct cga ggt tct aca atc tct ctg atg aga aca gac   480
Leu Asn Glu Glu Ser Arg Gly Ser Thr Ile Ser Leu Met Arg Thr Asp
145                 150                 155                 160

tgt cta taaaatccca cagtccttgg gaagttgggg accacataca ctgttgggat   536
Cys Leu

gtacattgat acaaccttta tgatggcaat ttgacaatat ttattaaaat aaaaaatggt 596

tattcccttc aaaaaaaaaa aaaaaaaaaa aaa                              629
```

<210> 20
<211> 162
<212> PRT
<213> Unknown

<400> 20

```
Asn Glu Leu Ile Pro Asn Leu Glu Lys Glu Asp Gly Ser Ile Leu Ile
  1               5                  10                  15

Cys Leu Tyr Glu Ser Tyr Phe Asp Pro Gly Lys Ser Ile Ser Glu Asn
```

20 25 30

Ile Val Ser Phe Ile Glu Lys Ser Tyr Lys Ser Ile Phe Val Leu Xaa
35 40 45

Pro Asn Phe Val Gln Asn Glu Trp Cys His Tyr Glu Phe Tyr Phe Ala
50 55 60

His His Asn Leu Phe His Glu Asn Ser Asp Xaa Ile Ile Leu Ile Leu
65 70 75 80

Leu Glu Pro Ile Pro Phe Tyr Cys Ile Pro Thr Arg Tyr His Lys Leu
85 90 95

Glu Ala Leu Leu Glu Lys Lys Ala Tyr Leu Glu Trp Pro Lys Asp Arg
100 105 110

Arg Lys Cys Gly Leu Phe Trp Ala Asn Leu Arg Ala Ala Val Asn Val
115 120 125

Asn Val Leu Ala Thr Arg Glu Met Tyr Glu Leu Gln Thr Phe Thr Glu
130 135 140

Leu Asn Glu Glu Ser Arg Gly Ser Thr Ile Ser Leu Met Arg Thr Asp
145 150 155 160

Cys Leu

<210> 21
<211> 427
<212> DNA
<213> Unknown

<220>
<223> Description of Unknown Organism:primate; surmised
Homo sapiens

<220>
<221> CDS
<222> (1)..(426)

<400> 21
aag aac tcc aaa gaa aac ctc cag ttt cat gct ttt att tca tat agt 48
Lys Asn Ser Lys Glu Asn Leu Gln Phe His Ala Phe Ile Ser Tyr Ser
1 5 10 15

gaa cat gat tct gcc tgg gtg aaa agt gaa ttg gta cct tac cta gaa 96
Glu His Asp Ser Ala Trp Val Lys Ser Glu Leu Val Pro Tyr Leu Glu
20 25 30

aaa gaa gat ata cag att tgt ctt cat gag aga aac ttt gtc cct ggc 144
Lys Glu Asp Ile Gln Ile Cys Leu His Glu Arg Asn Phe Val Pro Gly
35 40 45

aag agc att gtg gaa aat atc atc aac tgc att gag aag agt tac aag 192
Lys Ser Ile Val Glu Asn Ile Ile Asn Cys Ile Glu Lys Ser Tyr Lys

```
          50                  55                  60

     tcc atc ttt gtt ttg tct ccc aac ttt gtc cag agt gag tgg tgc cat   240
     Ser Ile Phe Val Leu Ser Pro Asn Phe Val Gln Ser Glu Trp Cys His
 5    65                  70                  75                  80

     tac gaa ctc tat ttt gcc cat cac aat ctc ttt cat gaa gga tct aat   288
     Tyr Glu Leu Tyr Phe Ala His His Asn Leu Phe His Glu Gly Ser Asn
                     85                  90                  95
10
     aac tta atc ctc atc tta ctg gaa ccc att cca cag aac agc att ccc   336
     Asn Leu Ile Leu Ile Leu Leu Glu Pro Ile Pro Gln Asn Ser Ile Pro
                100                 105                 110

15   aac aag tac cac aag ctg aag gct ctc atg acg cag cgg act tat ttg   384
     Asn Lys Tyr His Lys Leu Lys Ala Leu Met Thr Gln Arg Thr Tyr Leu
            115                 120                 125

     cag tgg ccc aag gag aaa agc aaa cgt ggg ctc ttt tgg gct a         427
20   Gln Trp Pro Lys Glu Lys Ser Lys Arg Gly Leu Phe Trp Ala
        130                 135                 140


     <210> 22
25   <211> 142
     <212> PRT
     <213> Unknown

     <400> 22
30   Lys Asn Ser Lys Glu Asn Leu Gln Phe His Ala Phe Ile Ser Tyr Ser
       1               5                  10                  15

     Glu His Asp Ser Ala Trp Val Lys Ser Glu Leu Val Pro Tyr Leu Glu
                  20                  25                  30
35
     Lys Glu Asp Ile Gln Ile Cys Leu His Glu Arg Asn Phe Val Pro Gly
              35                  40                  45

     Lys Ser Ile Val Glu Asn Ile Ile Asn Cys Ile Glu Lys Ser Tyr Lys
40        50                  55                  60

     Ser Ile Phe Val Leu Ser Pro Asn Phe Val Gln Ser Glu Trp Cys His
      65                  70                  75                  80

45   Tyr Glu Leu Tyr Phe Ala His His Asn Leu Phe His Glu Gly Ser Asn
                     85                  90                  95

     Asn Leu Ile Leu Ile Leu Leu Glu Pro Ile Pro Gln Asn Ser Ile Pro
                100                 105                 110
50
     Asn Lys Tyr His Lys Leu Lys Ala Leu Met Thr Gln Arg Thr Tyr Leu
            115                 120                 125

     Gln Trp Pro Lys Glu Lys Ser Lys Arg Gly Leu Phe Trp Ala
55      130                 135                 140


     <210> 23
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<211> 662
<212> DNA
<213> Unknown

<220>
<223> Description of Unknown Organism:primate; surmised Homo sapiens

<220>
<221> CDS
<222> (1)..(627)

<220>
<221> misc_feature
<222> (18)..(136)
<223> Xaa translation depends on genetic code

<400> 23

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gct tcc acc tgt gcc tgg cct ggc ttc cct ggc ggg ggc ggc aaa gtg   48
Ala Ser Thr Cys Ala Trp Pro Gly Phe Pro Gly Gly Gly Gly Lys Val
  1               5                  10                  15

ggc gar atg agg atg ccc tgc cct acg atg cct tcg tgg tct tcg aca   96
Gly Xaa Met Arg Met Pro Cys Pro Thr Met Pro Ser Trp Ser Ser Thr
             20                  25                  30

aaa cgc rga gcg cag tgg cag act ggg tgt aca acg agc ttc ggg ggc  144
Lys Arg Xaa Ala Gln Trp Gln Thr Gly Cys Thr Thr Ser Phe Gly Gly
          35                  40                  45

agc tgg agg agt gcc gtg ggc gct ggg cac tcc gcc tgt gcc tgg agg  192
Ser Trp Arg Ser Ala Val Gly Ala Gly His Ser Ala Cys Ala Trp Arg
      50                  55                  60

aac gcg act ggc tgc ctg gca aaa ccc tct ttg aga acc tgt ggg cct  240
Asn Ala Thr Gly Cys Leu Ala Lys Pro Ser Leu Arg Thr Cys Gly Pro
  65                  70                  75                  80

cgg tct atg gca gcc gca aga cgc tgt ttg tgc tgg ccc aca cgg acc  288
Arg Ser Met Ala Ala Ala Arg Arg Cys Leu Cys Trp Pro Thr Arg Thr
                  85                  90                  95

ggg tca gtg gtc tct tgc gcg cca ktt ntc ctg ctg gcc cag cag cgc  336
Gly Ser Val Val Ser Cys Ala Pro Xaa Xaa Leu Leu Ala Gln Gln Arg
             100                 105                 110

ctg ctg gar gac cgc aag gac gtc gtg gtg ctg gtg atc cta ang cct  384
Leu Leu Xaa Asp Arg Lys Asp Val Val Val Leu Val Ile Leu Xaa Pro
         115                 120                 125

gac ggc caa gcc tcc cga cta cnn gat gcg ctg acc agc gcc tct gcc  432
Asp Gly Gln Ala Ser Arg Leu Xaa Asp Ala Leu Thr Ser Ala Ser Ala
     130                 135                 140

gcc aga gtg tcc tcc tct ggc ccc acc agc cca gtg gtc gcg cag ctt  480
Ala Arg Val Ser Ser Ser Gly Pro Thr Ser Pro Val Val Ala Gln Leu
145                 150                 155                 160
```

```
ctg agg cca gca tgc atg gcc ctg acc agg gac aac cac cac ttc tat   528
Leu Arg Pro Ala Cys Met Ala Leu Thr Arg Asp Asn His His Phe Tyr
                165                 170                 175

aac cgg aac ttc tgc cag gga acc cac ggc cga ata gcc gtg agc cgg   576
Asn Arg Asn Phe Cys Gln Gly Thr His Gly Arg Ile Ala Val Ser Arg
            180                 185                 190

aat cct gca cgg tgc cac ctc cac aca cac cta aca tat gcc tgc ctg   624
Asn Pro Ala Arg Cys His Leu His Thr His Leu Thr Tyr Ala Cys Leu
        195                 200                 205

atc tgaccaacac atgctcgcca ccctcaccac acacc                       662
Ile
```

<210> 24
<211> 209
<212> PRT
<213> Unknown

<400> 24

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Ala Ser Thr Cys Ala Trp Pro Gly Phe Pro Gly Gly Gly Gly Lys Val
  1               5                  10                  15

Gly Xaa Met Arg Met Pro Cys Pro Thr Met Pro Ser Trp Ser Ser Thr
             20                  25                  30

Lys Arg Xaa Ala Gln Trp Gln Thr Gly Cys Thr Thr Ser Phe Gly Gly
         35                  40                  45

Ser Trp Arg Ser Ala Val Gly Ala Gly His Ser Ala Cys Ala Trp Arg
     50                  55                  60

Asn Ala Thr Gly Cys Leu Ala Lys Pro Ser Leu Arg Thr Cys Gly Pro
 65                  70                  75                  80

Arg Ser Met Ala Ala Ala Arg Arg Cys Leu Cys Trp Pro Thr Arg Thr
                 85                  90                  95

Gly Ser Val Val Ser Cys Ala Pro Xaa Xaa Leu Leu Ala Gln Gln Arg
            100                 105                 110

Leu Leu Xaa Asp Arg Lys Asp Val Val Val Leu Val Ile Leu Xaa Pro
        115                 120                 125

Asp Gly Gln Ala Ser Arg Leu Xaa Asp Ala Leu Thr Ser Ala Ser Ala
    130                 135                 140

Ala Arg Val Ser Ser Ser Gly Pro Thr Ser Pro Val Val Ala Gln Leu
145                 150                 155                 160

Leu Arg Pro Ala Cys Met Ala Leu Thr Arg Asp Asn His His Phe Tyr
                165                 170                 175

Asn Arg Asn Phe Cys Gln Gly Thr His Gly Arg Ile Ala Val Ser Arg
            180                 185                 190
```

```
Asn Pro Ala Arg Cys His Leu His Thr His Leu Thr Tyr Ala Cys Leu
        195                 200                 205

Ile

<210> 25
<211> 4865
<212> DNA
<213> Unknown

<220>
<223> Description of Unknown Organism:primate; surmised
      Homo sapiens

<220>
<221> CDS
<222> (107)..(2617)

<220>
<221> mat_peptide
<222> (173)..(2617)

<220>
<221> misc_feature
<222> (189)
<223> Xaa translation depends on genetic code

<400> 25
aaaatactcc cttgcctcaa aaactgctcg gtcaaacggt gatagcaaac cacgcattca 60

cagggccact gctgctcaca naascagtga ggatgatgcc aggatg atg tct gcc    115
                                                Met Ser Ala
                                                    -20

tcg cgc ctg gct ggg act ctg atc cca gcc atg gcc ttc ctc tcc tgc   163
Ser Arg Leu Ala Gly Thr Leu Ile Pro Ala Met Ala Phe Leu Ser Cys
                -15                 -10                 -5

gtg aga cca gaa agc tgg gag ccc tgc gtg gag gtt cct aat att act   211
Val Arg Pro Glu Ser Trp Glu Pro Cys Val Glu Val Pro Asn Ile Thr
         -1   1              5                  10

tat caa tgc atg gag ctg aat ttc tac aaa atc ccc gac aac ctc ccc   259
Tyr Gln Cys Met Glu Leu Asn Phe Tyr Lys Ile Pro Asp Asn Leu Pro
     15                  20                  25

ttc tca acc aag aac ctg gac ctg agc ttt aat ccc ctg agg cat tta   307
Phe Ser Thr Lys Asn Leu Asp Leu Ser Phe Asn Pro Leu Arg His Leu
 30                  35                  40                  45

ggc agc tat agc ttc ttc agt ttc cca gaa ctg cag gtg ctg gat tta   355
Gly Ser Tyr Ser Phe Phe Ser Phe Pro Glu Leu Gln Val Leu Asp Leu
                 50                  55                  60

tcc agg tgt gaa atc cag aca att gaa gat ggg gca tat cag agc cta   403
Ser Arg Cys Glu Ile Gln Thr Ile Glu Asp Gly Ala Tyr Gln Ser Leu
```

```
                65                  70                  75

     agc cac ctc tct acc tta ata ttg aca gga aac ccc atc cag agt tta   451
     Ser His Leu Ser Thr Leu Ile Leu Thr Gly Asn Pro Ile Gln Ser Leu
5              80                  85                  90

     gcc ctg gga gcc ttt tct gga cta tca agt tta cag aag ctg gtg gct   499
     Ala Leu Gly Ala Phe Ser Gly Leu Ser Ser Leu Gln Lys Leu Val Ala
          95                 100                 105
10
     gtg gag aca aat cta gca tct cta gag aac ttc ccc att gga cat ctc   547
     Val Glu Thr Asn Leu Ala Ser Leu Glu Asn Phe Pro Ile Gly His Leu
     110                 115                 120                 125

15   aaa act ttg aaa gaa ctt aat gtg gct cac aat ctt atc caa tct ttc   595
     Lys Thr Leu Lys Glu Leu Asn Val Ala His Asn Leu Ile Gln Ser Phe
                    130                 135                 140

     aaa tta cct gag tat ttt tct aat ctg acc aat cta gag cac ttg gac   643
20   Lys Leu Pro Glu Tyr Phe Ser Asn Leu Thr Asn Leu Glu His Leu Asp
                145                 150                 155

     ctt tcc agc aac aag att caa agt att tat tgc aca gac ttg cgg gtt   691
     Leu Ser Ser Asn Lys Ile Gln Ser Ile Tyr Cys Thr Asp Leu Arg Val
25             160                 165                 170

     cta cat caa atg ccc cta ctc aat ctc tct tta gac ctg tcc ctg aay   739
     Leu His Gln Met Pro Leu Leu Asn Leu Ser Leu Asp Leu Ser Leu Xaa
          175                 180                 185
30
     cct atg aac ttt atc caa cca ggt gca ttt aaa gaa att agg ctt cat   787
     Pro Met Asn Phe Ile Gln Pro Gly Ala Phe Lys Glu Ile Arg Leu His
     190                 195                 200                 205

35   aag ctg act tta aga aat aat ttt gat agt tta aat gta atg aaa act   835
     Lys Leu Thr Leu Arg Asn Asn Phe Asp Ser Leu Asn Val Met Lys Thr
                    210                 215                 220

     tgt att caa ggt ctg gct ggt tta gaa gtc cat cgt ttg gtt ctg gga   883
40   Cys Ile Gln Gly Leu Ala Gly Leu Glu Val His Arg Leu Val Leu Gly
                225                 230                 235

     gaa ttt aga aat gaa gga aac ttg gaa aag ttt gac aaa tct gct cta   931
     Glu Phe Arg Asn Glu Gly Asn Leu Glu Lys Phe Asp Lys Ser Ala Leu
45             240                 245                 250

     gag ggc ctg tgc aat ttg acc att gaa gaa ttc cga tta gca tac tta   979
     Glu Gly Leu Cys Asn Leu Thr Ile Glu Glu Phe Arg Leu Ala Tyr Leu
          255                 260                 265
50
     gac tac tac ctc gat gat att att gac tta ttt aat tgt ttg aca aat   1027
     Asp Tyr Tyr Leu Asp Asp Ile Ile Asp Leu Phe Asn Cys Leu Thr Asn
     270                 275                 280                 285

55   gtt tct tca ttt tcc ctg gtg agt gtg act att gaa agg gta aaa gac   1075
     Val Ser Ser Phe Ser Leu Val Ser Val Thr Ile Glu Arg Val Lys Asp
                    290                 295                 300
```

```
ttt tct tat aat ttc gga tgg caa cat tta gaa tta gtt aac tgt aaa   1123
Phe Ser Tyr Asn Phe Gly Trp Gln His Leu Glu Leu Val Asn Cys Lys
            305                 310                 315

ttt gga cag ttt ccc aca ttg aaa ctc aaa tct ctc aaa agg ctt act   1171
Phe Gly Gln Phe Pro Thr Leu Lys Leu Lys Ser Leu Lys Arg Leu Thr
        320                 325                 330

ttc act tcc aac aaa ggt ggg aat gct ttt tca gaa gtt gat cta cca   1219
Phe Thr Ser Asn Lys Gly Gly Asn Ala Phe Ser Glu Val Asp Leu Pro
    335                 340                 345

agc ctt gag ttt cta gat ctc agt aga aat ggc ttg agt ttc aaa ggt   1267
Ser Leu Glu Phe Leu Asp Leu Ser Arg Asn Gly Leu Ser Phe Lys Gly
350                 355                 360                 365

tgc tgt tct caa agt gat ttt ggg aca acc agc cta aag tat tta gat   1315
Cys Cys Ser Gln Ser Asp Phe Gly Thr Thr Ser Leu Lys Tyr Leu Asp
                370                 375                 380

ctg agc ttc aat ggt gtt att acc atg agt tca aac ttc ttg ggc tta   1363
Leu Ser Phe Asn Gly Val Ile Thr Met Ser Ser Asn Phe Leu Gly Leu
            385                 390                 395

gaa caa cta gaa cat ctg gat ttc cag cat tcc aat ttg aaa caa atg   1411
Glu Gln Leu Glu His Leu Asp Phe Gln His Ser Asn Leu Lys Gln Met
        400                 405                 410

agt gag ttt tca gta ttc cta tca ctc aga aac ctc att tac ctt gac   1459
Ser Glu Phe Ser Val Phe Leu Ser Leu Arg Asn Leu Ile Tyr Leu Asp
    415                 420                 425

att tct cat act cac acc aga gtt gct ttc aat ggc atc ttc aat ggc   1507
Ile Ser His Thr His Thr Arg Val Ala Phe Asn Gly Ile Phe Asn Gly
430                 435                 440                 445

ttg tcc agt ctc gaa gtc ttg aaa atg gct ggc aat tct ttc cag gaa   1555
Leu Ser Ser Leu Glu Val Leu Lys Met Ala Gly Asn Ser Phe Gln Glu
                450                 455                 460

aac ttc ctt cca gat atc ttc aca gag ctg aga aac ttg acc ttc ctg   1603
Asn Phe Leu Pro Asp Ile Phe Thr Glu Leu Arg Asn Leu Thr Phe Leu
            465                 470                 475

gac ctc tct cag tgt caa ctg gag cag ttg tct cca aca gca ttt aac   1651
Asp Leu Ser Gln Cys Gln Leu Glu Gln Leu Ser Pro Thr Ala Phe Asn
        480                 485                 490

tca ctc tcc agt ctt cag gta cta aat atg agc cac aac aac ttc ttt   1699
Ser Leu Ser Ser Leu Gln Val Leu Asn Met Ser His Asn Asn Phe Phe
    495                 500                 505

tca ttg gat acg ttt cct tat aag tgt ctg aac tcc ctc cag gtt ctt   1747
Ser Leu Asp Thr Phe Pro Tyr Lys Cys Leu Asn Ser Leu Gln Val Leu
510                 515                 520                 525

gat tac agt ctc aat cac ata atg act tcc aaa aaa cag gaa cta cag   1795
Asp Tyr Ser Leu Asn His Ile Met Thr Ser Lys Lys Gln Glu Leu Gln
```

```
                    530                 535                 540

     cat ttt cca agt agt cta gct ttc tta aat ctt act cag aat gac ttt   1843
     His Phe Pro Ser Ser Leu Ala Phe Leu Asn Leu Thr Gln Asn Asp Phe
 5               545                 550                 555

     gct tgt act tgt gaa cac cag agt ttc ctg caa tgg atc aag gac cag   1891
     Ala Cys Thr Cys Glu His Gln Ser Phe Leu Gln Trp Ile Lys Asp Gln
             560                 565                 570
10
     agg cag ctc ttg gtg gaa gtt gaa cga atg gaa tgt gca aca cct tca   1939
     Arg Gln Leu Leu Val Glu Val Glu Arg Met Glu Cys Ala Thr Pro Ser
         575                 580                 585

15   gat aag cag ggc atg cct gtg ctg agt ttg aat atc acc tgt cag atg   1987
     Asp Lys Gln Gly Met Pro Val Leu Ser Leu Asn Ile Thr Cys Gln Met
     590                 595                 600                 605

     aat aag acc atc att ggt gtg tcg gtc ctc agt gtg ctt gta gta tct   2035
20   Asn Lys Thr Ile Ile Gly Val Ser Val Leu Ser Val Leu Val Val Ser
                     610                 615                 620

     gtt gta gca gtt ctg gtc tat aag ttc tat ttt cac ctg atg ctt ctt   2083
     Val Val Ala Val Leu Val Tyr Lys Phe Tyr Phe His Leu Met Leu Leu
25               625                 630                 635

     gct ggc tgc ata aag tat ggt aga ggt gaa aac atc tat gat gcc ttt   2131
     Ala Gly Cys Ile Lys Tyr Gly Arg Gly Glu Asn Ile Tyr Asp Ala Phe
             640                 645                 650
30
     gtt atc tac tca agc cag gat gag gac tgg gta agg aat gag cta gta   2179
     Val Ile Tyr Ser Ser Gln Asp Glu Asp Trp Val Arg Asn Glu Leu Val
         655                 660                 665

35   aag aat tta gaa gaa ggg gtg cct cca ttt cag ctc tgc ctt cac tac   2227
     Lys Asn Leu Glu Glu Gly Val Pro Pro Phe Gln Leu Cys Leu His Tyr
     670                 675                 680                 685

     aga gac ttt att ccc ggt gtg gcc att gct gcc aac atc atc cat gaa   2275
40   Arg Asp Phe Ile Pro Gly Val Ala Ile Ala Ala Asn Ile Ile His Glu
                     690                 695                 700

     ggt ttc cat aaa agc cga aag gtg att gtt gtg gtg tcc cag cac ttc   2323
     Gly Phe His Lys Ser Arg Lys Val Ile Val Val Val Ser Gln His Phe
45               705                 710                 715

     atc cag agc cgc tgg tgt atc ttt gaa tat gag att gct cag acc tgg   2371
     Ile Gln Ser Arg Trp Cys Ile Phe Glu Tyr Glu Ile Ala Gln Thr Trp
             720                 725                 730
50
     cag ttt ctg agc agt cgt gct ggt atc atc ttc att gtc ctg cag aag   2419
     Gln Phe Leu Ser Ser Arg Ala Gly Ile Ile Phe Ile Val Leu Gln Lys
         735                 740                 745

55   gtg gag aag acc ctg ctc agg cag cag gtg gag ctg tac cgc ctt ctc   2467
     Val Glu Lys Thr Leu Leu Arg Gln Gln Val Glu Leu Tyr Arg Leu Leu
     750                 755                 760                 765
```

agc agg aac act tac ctg gag tgg gag gac agt gtc ctg ggg cgg cac 2515
Ser Arg Asn Thr Tyr Leu Glu Trp Glu Asp Ser Val Leu Gly Arg His
770 775 780

atc ttc tgg aga cga ctc aga aaa gcc ctg ctg gat ggt aaa tca tgg 2563
Ile Phe Trp Arg Arg Leu Arg Lys Ala Leu Leu Asp Gly Lys Ser Trp
785 790 795

aat cca gaa gga aca gtg ggt aca gga tgc aat tgg cag gaa gca aca 2611
Asn Pro Glu Gly Thr Val Gly Thr Gly Cys Asn Trp Gln Glu Ala Thr
800 805 810

tct atc tgaagaggaa aaataaaaac ctcctgaggc atttcttgcc cagctgggtc 2667
Ser Ile
815

caacacttgt tcagttaata agtattaaat gctgccacat gtcaggcctt atgctaaggg 2727

tgagtaattc catggtgcac tagatatgca gggctgctaa tctcaaggag cttccagtgc 2787

agagggaata aatgctagac taaaatacag agtcttccag gtgggcattt caaccaactc 2847

agtcaaggaa cccatgacaa agaaagtcat ttcaactctt acctcatcaa gttgaataaa 2907

gacagagaaa acagaaagag acattgttct tttcctgagt cttttgaatg gaaattgtat 2967

tatgttatag ccatcataaa accattttgg tagttttgac tgaactgggt gttcactttt 3027

tccttttttga ttgaatacaa tttaaattct acttgatgac tgcagtcgtc aaggggctcc 3087

tgatgcaaga tgccccttcc attttaagtc tgtctccttа cagakgttaa agtctantgg 3147

ctaattccta aggaaacctg attaacacat gctcacaacc atcctggtca ttctcganca 3207

tgttctattt tttaactaat caccccctgat atatttttat ttttatatat ccagttttca 3267

tttttttacg tcttgcctat aagctaatat cataaataag gttgtttaag acgtgcttca 3327

aatatccata ttaaccacta tttttcaagg aagtatggaa aagtacactc tgtcactttg 3387

tcactcgatg tcattccaaa gttattgcct actaagtaat gactgtcatg aaagcagcat 3447

tgaaataatt tgtttaaagg gggcactctt ttaaacggga agaaaatttc cgcttcctgg 3507

tcttatcatg gacaatttgg gctakaggca kgaaggaagt gggatkacct caggangtca 3567

ccttttcttg attccagaaa catatgggct gataaacccg gggtgacctc atgaaatgag 3627

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tagggagaca cagatggctg ggatccctcc cctgtaccct tctcactgmc aggagaacta 3747

cgtgtgaagg tattcaaggc agggagtata cattgctgtt tcctgttggg caatgctcct 3807

tgaccacatt ttgggaagag tggatgttat cattgagaaa acaatgtgtc tggaattaat 3867

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ggaacattca aggaaaagga caatcaggat gtcatcaggg aaatgaaaat aaaaaccaca 3987

atgagatatc accttatacc aggtagatgg ctactataaa aaaatgaagt gtcatcaagg 4047

atatagagaa attggaaccc ttcttcactg ctggagggaa tggaaaatgg tgtagccgtt 4107

atgaaaaaca gtacggaggt ttctcaaaaa ttaaaaatag aactgctata tgatccagca 4167

atctcacttc tgtatatata cccaaaataa ttgaaatcag aatttcaaga aaatatttac 4227

actcccatgt tcattgtggc actcttcaca atcactgttt ccaaagttat ggaaacaacc 4287

caaatttcca ttggaaaata aatggacaaa ggaaatgtgc atataacgta caatggggat 4347

attattcagc ctaaaaaaag gggggatcct gttatttatg acaacatgaa taaacccgga 4407

ggccattatg ctatgtaaaa tgagcaagta acagaaagac aaatactgcc tgatttcatt 4467

tatatgaggt tctaaaatag tcaaactcat agaagcagag aatagaacag tggttcctag 4527

ggaaaaggag gaagggagaa atgaggaaat agggagttgt ctaattggta taaaattata 4587

gtatgcaaga tgaattagct ctaaagatca gctgtatagc agagttcgta taatgaacaa 4647

tactgtatta tgcacttaac attttgttaa gagggtacct ctcatgttaa gtgttcttac 4707

catatacata tacacaagga agcttttgga ggtgatggat atatttatta ccttgattgt 4767

ggtgatggtt tgacaggtat gtgactatgt ctaaactcat caaattgtat acattaaata 4827

tatgcagttt tataatatca aaaaaaaaaa aaaaaaaa 4865

<210> 26
<211> 837
<212> PRT
<213> Unknown

<400> 26

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Met Ser Ala Ser Arg Leu Ala Gly Thr Leu Ile Pro Ala Met Ala Phe
        -20                 -15                 -10

Leu Ser Cys Val Arg Pro Glu Ser Trp Glu Pro Cys Val Glu Val Pro
     -5              -1   1               5                  10

Asn Ile Thr Tyr Gln Cys Met Glu Leu Asn Phe Tyr Lys Ile Pro Asp
                15                  20                  25

Asn Leu Pro Phe Ser Thr Lys Asn Leu Asp Leu Ser Phe Asn Pro Leu
             30                  35                  40

Arg His Leu Gly Ser Tyr Ser Phe Phe Ser Phe Pro Glu Leu Gln Val
         45                  50                  55

Leu Asp Leu Ser Arg Cys Glu Ile Gln Thr Ile Glu Asp Gly Ala Tyr
     60                  65                  70

Gln Ser Leu Ser His Leu Ser Thr Leu Ile Leu Thr Gly Asn Pro Ile
```

75 80 85 90

Gln Ser Leu Ala Leu Gly Ala Phe Ser Gly Leu Ser Ser Leu Gln Lys
95 100 105

Leu Val Ala Val Glu Thr Asn Leu Ala Ser Leu Glu Asn Phe Pro Ile
110 115 120

Gly His Leu Lys Thr Leu Lys Glu Leu Asn Val Ala His Asn Leu Ile
125 130 135

Gln Ser Phe Lys Leu Pro Glu Tyr Phe Ser Asn Leu Thr Asn Leu Glu
140 145 150

His Leu Asp Leu Ser Ser Asn Lys Ile Gln Ser Ile Tyr Cys Thr Asp
155 160 165 170

Leu Arg Val Leu His Gln Met Pro Leu Leu Asn Leu Ser Leu Asp Leu
175 180 185

Ser Leu Xaa Pro Met Asn Phe Ile Gln Pro Gly Ala Phe Lys Glu Ile
190 195 200

Arg Leu His Lys Leu Thr Leu Arg Asn Asn Phe Asp Ser Leu Asn Val
205 210 215

Met Lys Thr Cys Ile Gln Gly Leu Ala Gly Leu Glu Val His Arg Leu
220 225 230

Val Leu Gly Glu Phe Arg Asn Glu Gly Asn Leu Glu Lys Phe Asp Lys
235 240 245 250

Ser Ala Leu Glu Gly Leu Cys Asn Leu Thr Ile Glu Glu Phe Arg Leu
255 260 265

Ala Tyr Leu Asp Tyr Tyr Leu Asp Asp Ile Ile Asp Leu Phe Asn Cys
270 275 280

Leu Thr Asn Val Ser Ser Phe Ser Leu Val Ser Val Thr Ile Glu Arg
285 290 295

Val Lys Asp Phe Ser Tyr Asn Phe Gly Trp Gln His Leu Glu Leu Val
300 305 310

Asn Cys Lys Phe Gly Gln Phe Pro Thr Leu Lys Leu Lys Ser Leu Lys
315 320 325 330

Arg Leu Thr Phe Thr Ser Asn Lys Gly Gly Asn Ala Phe Ser Glu Val
335 340 345

Asp Leu Pro Ser Leu Glu Phe Leu Asp Leu Ser Arg Asn Gly Leu Ser
350 355 360

Phe Lys Gly Cys Cys Ser Gln Ser Asp Phe Gly Thr Thr Ser Leu Lys
365 370 375

Tyr Leu Asp Leu Ser Phe Asn Gly Val Ile Thr Met Ser Ser Asn Phe
380 385 390

```
Leu Gly Leu Glu Gln Leu Glu His Leu Asp Phe Gln His Ser Asn Leu
395                 400                 405                 410

Lys Gln Met Ser Glu Phe Ser Val Phe Leu Ser Leu Arg Asn Leu Ile
                415                 420                 425

Tyr Leu Asp Ile Ser His Thr His Thr Arg Val Ala Phe Asn Gly Ile
            430                 435                 440

Phe Asn Gly Leu Ser Ser Leu Glu Val Leu Lys Met Ala Gly Asn Ser
        445                 450                 455

Phe Gln Glu Asn Phe Leu Pro Asp Ile Phe Thr Glu Leu Arg Asn Leu
    460                 465                 470

Thr Phe Leu Asp Leu Ser Gln Cys Gln Leu Glu Gln Leu Ser Pro Thr
475                 480                 485                 490

Ala Phe Asn Ser Leu Ser Ser Leu Gln Val Leu Asn Met Ser His Asn
                495                 500                 505

Asn Phe Phe Ser Leu Asp Thr Phe Pro Tyr Lys Cys Leu Asn Ser Leu
            510                 515                 520

Gln Val Leu Asp Tyr Ser Leu Asn His Ile Met Thr Ser Lys Lys Gln
        525                 530                 535

Glu Leu Gln His Phe Pro Ser Ser Leu Ala Phe Leu Asn Leu Thr Gln
    540                 545                 550

Asn Asp Phe Ala Cys Thr Cys Glu His Gln Ser Phe Leu Gln Trp Ile
555                 560                 565                 570

Lys Asp Gln Arg Gln Leu Leu Val Glu Val Glu Arg Met Glu Cys Ala
                575                 580                 585

Thr Pro Ser Asp Lys Gln Gly Met Pro Val Leu Ser Leu Asn Ile Thr
            590                 595                 600

Cys Gln Met Asn Lys Thr Ile Ile Gly Val Ser Val Leu Ser Val Leu
        605                 610                 615

Val Val Ser Val Val Ala Val Leu Val Tyr Lys Phe Tyr Phe His Leu
    620                 625                 630

Met Leu Leu Ala Gly Cys Ile Lys Tyr Gly Arg Gly Glu Asn Ile Tyr
635                 640                 645                 650

Asp Ala Phe Val Ile Tyr Ser Ser Gln Asp Glu Asp Trp Val Arg Asn
                655                 660                 665

Glu Leu Val Lys Asn Leu Glu Glu Gly Val Pro Pro Phe Gln Leu Cys
            670                 675                 680

Leu His Tyr Arg Asp Phe Ile Pro Gly Val Ala Ile Ala Ala Asn Ile
        685                 690                 695
```

```
Ile His Glu Gly Phe His Lys Ser Arg Lys Val Ile Val Val Val Ser
    700                 705                 710

Gln His Phe Ile Gln Ser Arg Trp Cys Ile Phe Glu Tyr Glu Ile Ala
715                 720                 725                 730

Gln Thr Trp Gln Phe Leu Ser Ser Arg Ala Gly Ile Ile Phe Ile Val
                735                 740                 745

Leu Gln Lys Val Glu Lys Thr Leu Leu Arg Gln Gln Val Glu Leu Tyr
            750                 755                 760

Arg Leu Leu Ser Arg Asn Thr Tyr Leu Glu Trp Glu Asp Ser Val Leu
        765                 770                 775

Gly Arg His Ile Phe Trp Arg Arg Leu Arg Lys Ala Leu Leu Asp Gly
    780                 785                 790

Lys Ser Trp Asn Pro Glu Gly Thr Val Gly Thr Gly Cys Asn Trp Gln
795                 800                 805                 810

Glu Ala Thr Ser Ile
                815
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<210> 27
<211> 300
<212> DNA
<213> Unknown

<220>
<223> Description of Unknown Organism:rodent; surmised
Mus musculus

<220>
<221> CDS
<222> (1)..(300)

<220>
<221> misc_feature
<222> (62)..(100)
<223> Xaa translation depends on genetic code

<400> 27

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tcc tat tct atg gaa aaa gat gct ttc cta ttt atg aga aat ttg aag   48
Ser Tyr Ser Met Glu Lys Asp Ala Phe Leu Phe Met Arg Asn Leu Lys
  1               5                  10                  15

gtt ctc tca cta aaa gat aac aat gtc aca gct gtc ccc acc act ttg   96
Val Leu Ser Leu Lys Asp Asn Asn Val Thr Ala Val Pro Thr Thr Leu
             20                  25                  30

cca cct aat tta cta gag ctc tat ctt tat aac aat atc att aag aaa   144
Pro Pro Asn Leu Leu Glu Leu Tyr Leu Tyr Asn Asn Ile Ile Lys Lys
         35                  40                  45

atc caa gaa aat gat ttc aat aac ctc aat gag ttg caa gtn ctt gac   192
Ile Gln Glu Asn Asp Phe Asn Asn Leu Asn Glu Leu Gln Xaa Leu Asp
```

50 55 60

cta ngt gga aat tgc cct cga tgt nat aat gtc cca tat ccg tgt aca 240
Leu Xaa Gly Asn Cys Pro Arg Cys Xaa Asn Val Pro Tyr Pro Cys Thr
65 70 75 80

ccg tgt gaa aat aat tcc ccc tta cag atc cat gan aat gct ttc aat 288
Pro Cys Glu Asn Asn Ser Pro Leu Gln Ile His Xaa Asn Ala Phe Asn
85 90 95

tca tcg aca gan 300
Ser Ser Thr Xaa
100

<210> 28
<211> 100
<212> PRT
<213> Unknown

<400> 28
Ser Tyr Ser Met Glu Lys Asp Ala Phe Leu Phe Met Arg Asn Leu Lys
1 5 10 15

Val Leu Ser Leu Lys Asp Asn Asn Val Thr Ala Val Pro Thr Thr Leu
20 25 30

Pro Pro Asn Leu Leu Glu Leu Tyr Leu Tyr Asn Asn Ile Ile Lys Lys
35 40 45

Ile Gln Glu Asn Asp Phe Asn Asn Leu Asn Glu Leu Gln Xaa Leu Asp
50 55 60

Leu Xaa Gly Asn Cys Pro Arg Cys Xaa Asn Val Pro Tyr Pro Cys Thr
65 70 75 80

Pro Cys Glu Asn Asn Ser Pro Leu Gln Ile His Xaa Asn Ala Phe Asn
85 90 95

Ser Ser Thr Xaa
100

<210> 29
<211> 1756
<212> DNA
<213> Unknown

<220>
<223> Description of Unknown Organism:rodent; surmised
Mus musculus

<220>
<221> CDS
<222> (1)..(1182)

<400> 29
tct cca gaa att ccc tgg aat tcc ttg cct cct gag gtt ttt gag ggt 48

```
Ser Pro Glu Ile Pro Trp Asn Ser Leu Pro Pro Glu Val Phe Glu Gly
  1               5                  10                  15

atg ccg cca aat cta aag aat ctc tcc ttg gcc aaa aat ggg ctc aaa   96
Met Pro Pro Asn Leu Lys Asn Leu Ser Leu Ala Lys Asn Gly Leu Lys
            20                  25                  30

tct ttc ttt tgg gac aga ctc cag tta ctg aag cat ttg gaa att ttg  144
Ser Phe Phe Trp Asp Arg Leu Gln Leu Leu Lys His Leu Glu Ile Leu
        35                  40                  45

gac ctc agc cat aac cag ctg aca aaa gta cct gag aga ttg gcc aac  192
Asp Leu Ser His Asn Gln Leu Thr Lys Val Pro Glu Arg Leu Ala Asn
    50                  55                  60

tgt tcc aaa agt ctc aca aca ctg att ctt aag cat aat caa atc agg  240
Cys Ser Lys Ser Leu Thr Thr Leu Ile Leu Lys His Asn Gln Ile Arg
65                  70                  75                  80

caa ttg aca aaa tat ttt cta gaa gat gct ttg caa ttg cgc tat cta  288
Gln Leu Thr Lys Tyr Phe Leu Glu Asp Ala Leu Gln Leu Arg Tyr Leu
                85                  90                  95

gac atc agt tca aat aaa atc cag gtc att cag aag act agc ttc cca  336
Asp Ile Ser Ser Asn Lys Ile Gln Val Ile Gln Lys Thr Ser Phe Pro
            100                 105                 110

gaa aat gtc ctc aac aat ctg gag atg ttg gtt tta cat cac aat cgc  384
Glu Asn Val Leu Asn Asn Leu Glu Met Leu Val Leu His His Asn Arg
        115                 120                 125

ttt ctt tgc aac tgt gat gct gtg tgg ttt gtc tgg tgg gtt aac cat  432
Phe Leu Cys Asn Cys Asp Ala Val Trp Phe Val Trp Trp Val Asn His
    130                 135                 140

aca gat gtt act att cca tac ctg gcc act gat gtg act tgt gta ggt  480
Thr Asp Val Thr Ile Pro Tyr Leu Ala Thr Asp Val Thr Cys Val Gly
145                 150                 155                 160

cca gga gca cac aaa ggt caa agt gtc ata tcc ctt gat ctg tat acg  528
Pro Gly Ala His Lys Gly Gln Ser Val Ile Ser Leu Asp Leu Tyr Thr
                165                 170                 175

tgt gag tta gat ctc aca aac ctg att ctg ttc tca gtt tcc ata tca  576
Cys Glu Leu Asp Leu Thr Asn Leu Ile Leu Phe Ser Val Ser Ile Ser
            180                 185                 190

tca gtc ctc ttt ctt atg gta gtt atg aca aca agt cac ctc ttt ttc  624
Ser Val Leu Phe Leu Met Val Val Met Thr Thr Ser His Leu Phe Phe
        195                 200                 205

tgg gat atg tgg tac att tat tat ttt tgg aaa gca aag ata aag ggg  672
Trp Asp Met Trp Tyr Ile Tyr Tyr Phe Trp Lys Ala Lys Ile Lys Gly
    210                 215                 220

tat cca gca tct gca atc cca tgg agt cct tgt tat gat gct ttt att  720
Tyr Pro Ala Ser Ala Ile Pro Trp Ser Pro Cys Tyr Asp Ala Phe Ile
225                 230                 235                 240
```

```
gtg tat gac act aaa aac tca gct gtg aca gaa tgg gtt ttg cag gag   768
Val Tyr Asp Thr Lys Asn Ser Ala Val Thr Glu Trp Val Leu Gln Glu
                245                 250                 255

ctg gtg gca aaa ttg gaa gat cca aga gaa aaa cac ttc aat ttg tgt   816
Leu Val Ala Lys Leu Glu Asp Pro Arg Glu Lys His Phe Asn Leu Cys
            260                 265                 270

cta gaa gaa aga gac tgg cta cca gga cag cca gtt cta gaa aac ctt   864
Leu Glu Glu Arg Asp Trp Leu Pro Gly Gln Pro Val Leu Glu Asn Leu
        275                 280                 285

tcc cag agc ata cag ctc agc aaa aag aca gtg ttt gtg atg aca cag   912
Ser Gln Ser Ile Gln Leu Ser Lys Lys Thr Val Phe Val Met Thr Gln
    290                 295                 300

aaa tat gct aag act gag agt ttt aag atg gca ttt tat ttg tct cat   960
Lys Tyr Ala Lys Thr Glu Ser Phe Lys Met Ala Phe Tyr Leu Ser His
305                 310                 315                 320

cag agg ctc ctg gat gaa aaa gtg gat gtg att atc ttg ata ttc ttg  1008
Gln Arg Leu Leu Asp Glu Lys Val Asp Val Ile Ile Leu Ile Phe Leu
                325                 330                 335

gaa aga cct ctt cag aag tct aag ttt ctt cag ctc agg aag aga ctc  1056
Glu Arg Pro Leu Gln Lys Ser Lys Phe Leu Gln Leu Arg Lys Arg Leu
            340                 345                 350

tgc agg agc tct gtc ctt gag tgg cct gca aat cca cag gct cac cca  1104
Cys Arg Ser Ser Val Leu Glu Trp Pro Ala Asn Pro Gln Ala His Pro
        355                 360                 365

tac ttc tgg cag tgc ctg aaa aat gcc ctg acc aca gac aat cat gtg  1152
Tyr Phe Trp Gln Cys Leu Lys Asn Ala Leu Thr Thr Asp Asn His Val
    370                 375                 380

gct tat agt caa atg ttc aag gaa aca gtc tagctctctg aagaatgtca    1202
Ala Tyr Ser Gln Met Phe Lys Glu Thr Val
385                 390

ccacctagga catgccttgg tacctgaagt tttcataaag gtttccataa atgaaggtct 1262

gaatttttcc taacagttgt catggctcag attggtggga aatcatcaat atatggctaa 1322

gaaattaaga aggggagact gatagaagat aatttctttc ttcatgtgcc atgctcagtt 1382

aaatatttcc cctagctcaa atctgaaaaa ctgtgcctag gagacaacac aaggctttga 1442

tttatctgca tacaattgat aagagccaca catctgcccc gaagaagtac tagtagtttt 1502

agtagtaggg taaaaattac acaagctttc tctctctctg atactgaact gtaccagagt 1562

tcaatgaaat aaaagcccag agaacttctc agtaaatggt ttcattatca tgtagtatcc 1622

accatgcaat atgccacaaa rccgctactg gtacaggaca gntggtagct gcttcaakgc 1682

ctcttatcat tttcttgggg cccatggagg ggttctytgg gaaadaggga agkttttttt 1742
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tggccatcca tgaa 1756

<210> 30
<211> 394
<212> PRT
<213> Unknown

<400> 30

Ser Pro Glu Ile Pro Trp Asn Ser Leu Pro Pro Glu Val Phe Glu Gly
1 5 10 15

Met Pro Pro Asn Leu Lys Asn Leu Ser Leu Ala Lys Asn Gly Leu Lys
20 25 30

Ser Phe Phe Trp Asp Arg Leu Gln Leu Leu Lys His Leu Glu Ile Leu
35 40 45

Asp Leu Ser His Asn Gln Leu Thr Lys Val Pro Glu Arg Leu Ala Asn
50 55 60

Cys Ser Lys Ser Leu Thr Thr Leu Ile Leu Lys His Asn Gln Ile Arg
65 70 75 80

Gln Leu Thr Lys Tyr Phe Leu Glu Asp Ala Leu Gln Leu Arg Tyr Leu
85 90 95

Asp Ile Ser Ser Asn Lys Ile Gln Val Ile Gln Lys Thr Ser Phe Pro
100 105 110

Glu Asn Val Leu Asn Asn Leu Glu Met Leu Val Leu His His Asn Arg
115 120 125

Phe Leu Cys Asn Cys Asp Ala Val Trp Phe Val Trp Trp Val Asn His
130 135 140

Thr Asp Val Thr Ile Pro Tyr Leu Ala Thr Asp Val Thr Cys Val Gly
145 150 155 160

Pro Gly Ala His Lys Gly Gln Ser Val Ile Ser Leu Asp Leu Tyr Thr
165 170 175

Cys Glu Leu Asp Leu Thr Asn Leu Ile Leu Phe Ser Val Ser Ile Ser
180 185 190

Ser Val Leu Phe Leu Met Val Val Met Thr Thr Ser His Leu Phe Phe
195 200 205

Trp Asp Met Trp Tyr Ile Tyr Tyr Phe Trp Lys Ala Lys Ile Lys Gly
210 215 220

Tyr Pro Ala Ser Ala Ile Pro Trp Ser Pro Cys Tyr Asp Ala Phe Ile
225 230 235 240

Val Tyr Asp Thr Lys Asn Ser Ala Val Thr Glu Trp Val Leu Gln Glu
245 250 255

```
Leu Val Ala Lys Leu Glu Asp Pro Arg Glu Lys His Phe Asn Leu Cys
            260                 265                 270

Leu Glu Glu Arg Asp Trp Leu Pro Gly Gln Pro Val Leu Glu Asn Leu
        275                 280                 285

Ser Gln Ser Ile Gln Leu Ser Lys Lys Thr Val Phe Val Met Thr Gln
    290                 295                 300

Lys Tyr Ala Lys Thr Glu Ser Phe Lys Met Ala Phe Tyr Leu Ser His
305                 310                 315                 320

Gln Arg Leu Leu Asp Glu Lys Val Asp Val Ile Ile Leu Ile Phe Leu
                325                 330                 335

Glu Arg Pro Leu Gln Lys Ser Lys Phe Leu Gln Leu Arg Lys Arg Leu
            340                 345                 350

Cys Arg Ser Ser Val Leu Glu Trp Pro Ala Asn Pro Gln Ala His Pro
        355                 360                 365

Tyr Phe Trp Gln Cys Leu Lys Asn Ala Leu Thr Thr Asp Asn His Val
    370                 375                 380

Ala Tyr Ser Gln Met Phe Lys Glu Thr Val
385                 390


<210> 31
<211> 999
<212> DNA
<213> Unknown

<220>
<223> Description of Unknown Organism:primate; surmised
      Homo sapiens

<220>
<221> CDS
<222> (2)..(847)

<220>
<221> misc_feature
<222> (1)..(282)
<223> Xaa translation depends on genetic code

<400> 31
c tcn gat gcc aag att cgg cac nag gca tat tca gag gtc atg atg gtt 49
  Xaa Asp Ala Lys Ile Arg His Xaa Ala Tyr Ser Glu Val Met Met Val
    1               5                  10                  15

gga tgg tca gat tca tac acc tgt gaa tac cct tta aac cta agg gga   97
Gly Trp Ser Asp Ser Tyr Thr Cys Glu Tyr Pro Leu Asn Leu Arg Gly
            20                  25                  30

act agg tta aaa gac gtt cat ctc cac gaa tta tct tgc aac aca gct  145
Thr Arg Leu Lys Asp Val His Leu His Glu Leu Ser Cys Asn Thr Ala
        35                  40                  45
```

```
ctg ttg att gtc acc att gtg gtt att atg cta gtt ctg ggg ttg gct   193
Leu Leu Ile Val Thr Ile Val Val Ile Met Leu Val Leu Gly Leu Ala
     50                  55                  60

gtg gcc ttc tgc tgt ctc cac ttt gat ctg ccc tgg tat ctc agg atg   241
Val Ala Phe Cys Cys Leu His Phe Asp Leu Pro Trp Tyr Leu Arg Met
 65                  70                  75                  80

cta ggt caa tgc aca caa aca tgg cac agg gtt agg aaa aca acc caa   289
Leu Gly Gln Cys Thr Gln Thr Trp His Arg Val Arg Lys Thr Thr Gln
                85                  90                  95

gaa caa ctc aag aga aat gtc cga ttc cac gca ttt att tca tac agt   337
Glu Gln Leu Lys Arg Asn Val Arg Phe His Ala Phe Ile Ser Tyr Ser
            100                 105                 110

gaa cat gat tct ctg tgg gtg aag aat gaa ttg atc ccc aat cta gag   385
Glu His Asp Ser Leu Trp Val Lys Asn Glu Leu Ile Pro Asn Leu Glu
        115                 120                 125

aag gaa gat ggt tct atc ttg att tgc ctt tat gaa agc tac ttt gac   433
Lys Glu Asp Gly Ser Ile Leu Ile Cys Leu Tyr Glu Ser Tyr Phe Asp
    130                 135                 140

cct ggc aaa agc att agt gaa aat att gta agc ttc att gag aaa agc   481
Pro Gly Lys Ser Ile Ser Glu Asn Ile Val Ser Phe Ile Glu Lys Ser
145                 150                 155                 160

tat aag tcc atc ttt gtt ttg tct ccc aac ttt gtc cag aat gag tgg   529
Tyr Lys Ser Ile Phe Val Leu Ser Pro Asn Phe Val Gln Asn Glu Trp
                165                 170                 175

tgc cat tat gaa ttc tac ttt gcc cac cac aat ctc ttc cat gaa aat   577
Cys His Tyr Glu Phe Tyr Phe Ala His His Asn Leu Phe His Glu Asn
            180                 185                 190

tct gat cac ata att ctt atc tta ctg gaa ccc att cca ttc tat tgc   625
Ser Asp His Ile Ile Leu Ile Leu Leu Glu Pro Ile Pro Phe Tyr Cys
        195                 200                 205

att ccc acc agg tat cat aaa ctg raa gct ctc ctg gaa aaa aaa gca   673
Ile Pro Thr Arg Tyr His Lys Leu Xaa Ala Leu Leu Glu Lys Lys Ala
    210                 215                 220

tac ttg gaa tgg ccc aag gat agg cgt aaa tgt ggg ctt tty tgg gca   721
Tyr Leu Glu Trp Pro Lys Asp Arg Arg Lys Cys Gly Leu Xaa Trp Ala
225                 230                 235                 240

aac ctt cga gct gct gtt aat gtt aat gta tta gcc acc aga gaa atg   769
Asn Leu Arg Ala Ala Val Asn Val Asn Val Leu Ala Thr Arg Glu Met
                245                 250                 255

tat gaa ctg cag aca ttc aca gag tta aat gaa gag tct cga ggt tct   817
Tyr Glu Leu Gln Thr Phe Thr Glu Leu Asn Glu Glu Ser Arg Gly Ser
            260                 265                 270

aca atc tyt ctg atg aga aca gac tgt yta taaaatccca cagtccttgg     867
```

Thr Ile Xaa Leu Met Arg Thr Asp Cys Xaa
275 280

gaagttgggg accacataca ctgttgggat gtacattgat acaaccttta tgatggcaat 927

ttgacaatat ttattaaaat aaaaaatggt tattcccttc aaaaaaaaaa aaaaaaaaaa 987

aaaaaaaaaa aa 999

<210> 32
<211> 282
<212> PRT
<213> Unknown

<400> 32
Xaa Asp Ala Lys Ile Arg His Xaa Ala Tyr Ser Glu Val Met Met Val
1 5 10 15

Gly Trp Ser Asp Ser Tyr Thr Cys Glu Tyr Pro Leu Asn Leu Arg Gly
20 25 30

Thr Arg Leu Lys Asp Val His Leu His Glu Leu Ser Cys Asn Thr Ala
35 40 45

Leu Leu Ile Val Thr Ile Val Val Ile Met Leu Val Leu Gly Leu Ala
50 55 60

Val Ala Phe Cys Cys Leu His Phe Asp Leu Pro Trp Tyr Leu Arg Met
65 70 75 80

Leu Gly Gln Cys Thr Gln Thr Trp His Arg Val Arg Lys Thr Thr Gln
85 90 95

Glu Gln Leu Lys Arg Asn Val Arg Phe His Ala Phe Ile Ser Tyr Ser
100 105 110

Glu His Asp Ser Leu Trp Val Lys Asn Glu Leu Ile Pro Asn Leu Glu
115 120 125

Lys Glu Asp Gly Ser Ile Leu Ile Cys Leu Tyr Glu Ser Tyr Phe Asp
130 135 140

Pro Gly Lys Ser Ile Ser Glu Asn Ile Val Ser Phe Ile Glu Lys Ser
145 150 155 160

Tyr Lys Ser Ile Phe Val Leu Ser Pro Asn Phe Val Gln Asn Glu Trp
165 170 175

Cys His Tyr Glu Phe Tyr Phe Ala His His Asn Leu Phe His Glu Asn
180 185 190

Ser Asp His Ile Ile Leu Ile Leu Leu Glu Pro Ile Pro Phe Tyr Cys
195 200 205

Ile Pro Thr Arg Tyr His Lys Leu Xaa Ala Leu Leu Glu Lys Lys Ala
210 215 220

```
Tyr Leu Glu Trp Pro Lys Asp Arg Arg Lys Cys Gly Leu Xaa Trp Ala
225                 230                 235                 240

Asn Leu Arg Ala Ala Val Asn Val Asn Val Leu Ala Thr Arg Glu Met
                245                 250                 255

Tyr Glu Leu Gln Thr Phe Thr Glu Leu Asn Glu Glu Ser Arg Gly Ser
            260                 265                 270

Thr Ile Xaa Leu Met Arg Thr Asp Cys Xaa
        275                 280
```

<210> 33
<211> 1173
<212> DNA
<213> Unknown

<220>
<223> Description of Unknown Organism:primate; surmised Homo sapiens

<220>
<221> CDS
<222> (1)..(1008)

<220>
<221> misc_feature
<222> (285)
<223> Xaa translation depends on genetic code

<400> 33

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ctg cct gct ggc acc cgg ctc cgg agg ctg gat gtc agc tgc aac agc   48
Leu Pro Ala Gly Thr Arg Leu Arg Arg Leu Asp Val Ser Cys Asn Ser
 1               5                  10                  15

atc agc ttc gtg gcc ccc ggc ttc ttt tcc aag gcc aag gag ctg cga   96
Ile Ser Phe Val Ala Pro Gly Phe Phe Ser Lys Ala Lys Glu Leu Arg
            20                  25                  30

gag ctc aac ctt agc gcc aac gcc ctc aag aca gtg gac cac tcc tgg  144
Glu Leu Asn Leu Ser Ala Asn Ala Leu Lys Thr Val Asp His Ser Trp
        35                  40                  45

ttt ggg ccc ctg gcg agt gcc ctg caa ata cta gat gta agc gcc aac  192
Phe Gly Pro Leu Ala Ser Ala Leu Gln Ile Leu Asp Val Ser Ala Asn
    50                  55                  60

cct ctg cac tgc gcc tgt ggg gcg gcc ttt atg gac ttc ctg ctg gag  240
Pro Leu His Cys Ala Cys Gly Ala Ala Phe Met Asp Phe Leu Leu Glu
65                  70                  75                  80

gtg cag gct gcc gtg ccc ggt ctg ccc agc cgg gtg aag tgt ggc agt  288
Val Gln Ala Ala Val Pro Gly Leu Pro Ser Arg Val Lys Cys Gly Ser
                85                  90                  95

ccg ggc cag ctc cag ggc ctc agc atc ttt gca cag gac ctg cgc ctc  336
Pro Gly Gln Leu Gln Gly Leu Ser Ile Phe Ala Gln Asp Leu Arg Leu
```

```
                100                 105                 110

    tgc ctg gat gag gcc ctc tcc tgg gac tgt ttc gcc ctc tcg ctg ctg   384
    Cys Leu Asp Glu Ala Leu Ser Trp Asp Cys Phe Ala Leu Ser Leu Leu
5           115                 120                 125

    gct gtg gct ctg ggc ctg ggt gtg ccc atg ctg cat cac ctc tgt ggc   432
    Ala Val Ala Leu Gly Leu Gly Val Pro Met Leu His His Leu Cys Gly
        130                 135                 140
10
    tgg gac ctc tgg tac tgc ttc cac ctg tgc ctg gcc tgg ctt ccc tgg   480
    Trp Asp Leu Trp Tyr Cys Phe His Leu Cys Leu Ala Trp Leu Pro Trp
    145                 150                 155                 160

15  cgg ggg cgg caa agt ggg cga gat gag gat gcc ctg ccc tac gat gcc   528
    Arg Gly Arg Gln Ser Gly Arg Asp Glu Asp Ala Leu Pro Tyr Asp Ala
                    165                 170                 175

    ttc gtg gtc ttc gac aaa acg cag agc gca gtg gca gac tgg gtg tac   576
20  Phe Val Val Phe Asp Lys Thr Gln Ser Ala Val Ala Asp Trp Val Tyr
                180                 185                 190

    aac gag ctt cgg ggg cag ctg gag gag tgc cgt ggg cgc tgg gca ctc   624
    Asn Glu Leu Arg Gly Gln Leu Glu Glu Cys Arg Gly Arg Trp Ala Leu
25          195                 200                 205

    cgc ctg tgc ctg gag gaa cgc gac tgg ctg cct ggc aaa acc ctc ttt   672
    Arg Leu Cys Leu Glu Glu Arg Asp Trp Leu Pro Gly Lys Thr Leu Phe
        210                 215                 220
30
    gag aac ctg tgg gcc tcg gtc tat ggc agc cgc aag acg ctg ttt gtg   720
    Glu Asn Leu Trp Ala Ser Val Tyr Gly Ser Arg Lys Thr Leu Phe Val
    225                 230                 235                 240

35  ctg gcc cac acg gac cgg gtc agt ggt ctc ttg cgc gcc agc ttc ctg   768
    Leu Ala His Thr Asp Arg Val Ser Gly Leu Leu Arg Ala Ser Phe Leu
                    245                 250                 255

    ctg gcc cag cag cgc ctg ctg gag gac cgc aag gac gtc gtg gtg ctg   816
40  Leu Ala Gln Gln Arg Leu Leu Glu Asp Arg Lys Asp Val Val Val Leu
                260                 265                 270

    gtg atc ctg agc cct gac ggc cgc cgc tcc cgc tac gkg cgg ctg cgc   864
    Val Ile Leu Ser Pro Asp Gly Arg Arg Ser Arg Tyr Xaa Arg Leu Arg
45          275                 280                 285

    cag cgc ctc tgc cgc cag agt gtc ctc ctc tgg ccc cac cag ccc agt   912
    Gln Arg Leu Cys Arg Gln Ser Val Leu Leu Trp Pro His Gln Pro Ser
        290                 295                 300
50
    ggt cag cgc agc ttc tgg gcc cag ctg ggc atg gcc ctg acc agg gac   960
    Gly Gln Arg Ser Phe Trp Ala Gln Leu Gly Met Ala Leu Thr Arg Asp
    305                 310                 315                 320

55  aac cac cac ttc tat aac cgg aac ttc tgc cag gga ccc acg gcc gaa  1008
    Asn His His Phe Tyr Asn Arg Asn Phe Cys Gln Gly Pro Thr Ala Glu
                    325                 330                 335
```

tagccgtgag ccggaatcct gcacggtgcc acctccacac tcacctcacc tctgcctgcc 1068

tggtctgacc ctcccctgct cgcctcccte accccacacc tgacacagag caggcactca 1128

ataaatgcta ccgaaggcta aaaaaaaaaa aaaaaaaaaa aanna 1173

<210> 34
<211> 336
<212> PRT
<213> Unknown

<400> 34

Leu	Pro	Ala	Gly	Thr	Arg	Leu	Arg	Arg	Leu	Asp	Val	Ser	Cys	Asn	Ser
1				5				10					15		
Ile	Ser	Phe	Val	Ala	Pro	Gly	Phe	Phe	Ser	Lys	Ala	Lys	Glu	Leu	Arg
			20					25					30		
Glu	Leu	Asn	Leu	Ser	Ala	Asn	Ala	Leu	Lys	Thr	Val	Asp	His	Ser	Trp
		35					40					45			
Phe	Gly	Pro	Leu	Ala	Ser	Ala	Leu	Gln	Ile	Leu	Asp	Val	Ser	Ala	Asn
	50					55					60				
Pro	Leu	His	Cys	Ala	Cys	Gly	Ala	Ala	Phe	Met	Asp	Phe	Leu	Leu	Glu
65					70					75					80
Val	Gln	Ala	Ala	Val	Pro	Gly	Leu	Pro	Ser	Arg	Val	Lys	Cys	Gly	Ser
			85					90					95		
Pro	Gly	Gln	Leu	Gln	Gly	Leu	Ser	Ile	Phe	Ala	Gln	Asp	Leu	Arg	Leu
		100					105					110			
Cys	Leu	Asp	Glu	Ala	Leu	Ser	Trp	Asp	Cys	Phe	Ala	Leu	Ser	Leu	Leu
	115					120					125				
Ala	Val	Ala	Leu	Gly	Leu	Gly	Val	Pro	Met	Leu	His	His	Leu	Cys	Gly
130					135					140					
Trp	Asp	Leu	Trp	Tyr	Cys	Phe	His	Leu	Cys	Leu	Ala	Trp	Leu	Pro	Trp
145				150					155						160
Arg	Gly	Arg	Gln	Ser	Gly	Arg	Asp	Glu	Asp	Ala	Leu	Pro	Tyr	Asp	Ala
			165					170					175		
Phe	Val	Val	Phe	Asp	Lys	Thr	Gln	Ser	Ala	Val	Ala	Asp	Trp	Val	Tyr
		180					185					190			
Asn	Glu	Leu	Arg	Gly	Gln	Leu	Glu	Glu	Cys	Arg	Gly	Arg	Trp	Ala	Leu
	195					200					205				
Arg	Leu	Cys	Leu	Glu	Glu	Arg	Asp	Trp	Leu	Pro	Gly	Lys	Thr	Leu	Phe
210					215					220					
Glu	Asn	Leu	Trp	Ala	Ser	Val	Tyr	Gly	Ser	Arg	Lys	Thr	Leu	Phe	Val
225				230					235						240

Leu Ala His Thr Asp Arg Val Ser Gly Leu Leu Arg Ala Ser Phe Leu
245 250 255

Leu Ala Gln Gln Arg Leu Leu Glu Asp Arg Lys Asp Val Val Val Leu
260 265 270

Val Ile Leu Ser Pro Asp Gly Arg Arg Ser Arg Tyr Xaa Arg Leu Arg
275 280 285

Gln Arg Leu Cys Arg Gln Ser Val Leu Leu Trp Pro His Gln Pro Ser
290 295 300

Gly Gln Arg Ser Phe Trp Ala Gln Leu Gly Met Ala Leu Thr Arg Asp
305 310 315 320

Asn His His Phe Tyr Asn Arg Asn Phe Cys Gln Gly Pro Thr Ala Glu
325 330 335

<210> 35
<211> 497
<212> DNA
<213> Unknown

<220>
<223> Description of Unknown Organism:rodent; surmised
Mus musculus

<400> 35
tggcccacac ggaccgcgtc agtggcctcc tgcgcaccag cttcctgctg gctcagcagc 60

gcctgttgga agaccgcaag gacgtggtgg tgttggtgat cctgcgtccg gatgccccac 120

cgtcccgcta tgtgcgactg cgccagcgtc tctgccgcca gagtgtgctc ttctggcccc 180

agcgacccaa cgggcagggg ggcttctggg cccagctgag tacagccctg actagggaca 240

accgccactt ctataaccag aacttctgcc ggggacctac agcagaatag ctcagagcaa 300

cagctggaaa cagctgcatc ttcatgtctg gttcccgagt tgctctgcct gccttgctct 360

gtcttactac accgctattt ggcaagtgcg caatatatgc taccaagcca ccaggcccac 420

ggagcaaagg ttggctgtaa agggtagttt tcttcccatg catctttcag gagagtgaag 480

atagacacca aacccac 497

<210> 36
<211> 3099
<212> DNA
<213> Unknown

<220>
<223> Description of Unknown Organism:primate; surmised
Homo sapiens

<220>

<221> CDS
<222> (1)..(3096)

<220>
<221> mat_peptide
<222> (52)..(3096)

<220>
<221> misc_feature
<222> (725)
<223> Xaa translation depends on genetic code

<400> 36

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atg ctg acc tgc att ttc ctg cta ata tct ggt tcc tgt gag tta tgc   48
Met Leu Thr Cys Ile Phe Leu Leu Ile Ser Gly Ser Cys Glu Leu Cys
        -15                 -10                 -5

gcc gaa gaa aat ttt tct aga agc tat cct tgt gat gag aaa aag caa   96
Ala Glu Glu Asn Phe Ser Arg Ser Tyr Pro Cys Asp Glu Lys Lys Gln
 -1   1               5                  10                  15

aat gac tca gtt att gca gag tgc agc aat cgt cga cta cag gaa gtt   144
Asn Asp Ser Val Ile Ala Glu Cys Ser Asn Arg Arg Leu Gln Glu Val
                20                  25                  30

ccc caa acg gtg ggc aaa tat gtg aca gaa cta gac ctg tct gat aat   192
Pro Gln Thr Val Gly Lys Tyr Val Thr Glu Leu Asp Leu Ser Asp Asn
            35                  40                  45

ttc atc aca cac ata acg aat gaa tca ttt caa ggg ctg caa aat ctc   240
Phe Ile Thr His Ile Thr Asn Glu Ser Phe Gln Gly Leu Gln Asn Leu
        50                  55                  60

act aaa ata aat cta aac cac aac ccc aat gta cag cac cag aac gga   288
Thr Lys Ile Asn Leu Asn His Asn Pro Asn Val Gln His Gln Asn Gly
    65                  70                  75

aat ccc ggt ata caa tca aat ggc ttg aat atc aca gac ggg gca ttc   336
Asn Pro Gly Ile Gln Ser Asn Gly Leu Asn Ile Thr Asp Gly Ala Phe
80                  85                  90                  95

ctc aac cta aaa aac cta agg gag tta ctg ctt gaa gac aac cag tta   384
Leu Asn Leu Lys Asn Leu Arg Glu Leu Leu Leu Glu Asp Asn Gln Leu
                100                 105                 110

ccc caa ata ccc tct ggt ttg cca gag tct ttg aca gaa ctt agt cta   432
Pro Gln Ile Pro Ser Gly Leu Pro Glu Ser Leu Thr Glu Leu Ser Leu
            115                 120                 125

att caa aac aat ata tac aac ata act aaa gag ggc att tca aga ctt   480
Ile Gln Asn Asn Ile Tyr Asn Ile Thr Lys Glu Gly Ile Ser Arg Leu
        130                 135                 140

ata aac ttg aaa aat ctc tat ttg gcc tgg aac tgc tat ttt aac aaa   528
Ile Asn Leu Lys Asn Leu Tyr Leu Ala Trp Asn Cys Tyr Phe Asn Lys
    145                 150                 155

gtt tgc gag aaa act aac ata gaa gat gga gta ttt gaa acg ctg aca   576
```

```
Val Cys Glu Lys Thr Asn Ile Glu Asp Gly Val Phe Glu Thr Leu Thr
160                 165                 170                 175

aat ttg gag ttg cta tca cta tct ttc aat tct ctt tca cat gtg cca   624
Asn Leu Glu Leu Leu Ser Leu Ser Phe Asn Ser Leu Ser His Val Pro
                180                 185                 190

ccc aaa ctg cca agc tcc cta cgc aaa ctt ttt ctg agc aac acc cag   672
Pro Lys Leu Pro Ser Ser Leu Arg Lys Leu Phe Leu Ser Asn Thr Gln
            195                 200                 205

atc aaa tac att agt gaa gaa gat ttc aag gga ttg ata aat tta aca   720
Ile Lys Tyr Ile Ser Glu Glu Asp Phe Lys Gly Leu Ile Asn Leu Thr
        210                 215                 220

tta cta gat tta agc ggg aac tgt ccg agg tgc ttc aat gcc cca ttt   768
Leu Leu Asp Leu Ser Gly Asn Cys Pro Arg Cys Phe Asn Ala Pro Phe
    225                 230                 235

cca tgc gtg cct tgt gat ggt ggt gct tca att aat ata gat cgt ttt   816
Pro Cys Val Pro Cys Asp Gly Gly Ala Ser Ile Asn Ile Asp Arg Phe
240                 245                 250                 255

gct ttt caa aac ttg acc caa ctt cga tac cta aac ctc tct agc act   864
Ala Phe Gln Asn Leu Thr Gln Leu Arg Tyr Leu Asn Leu Ser Ser Thr
                260                 265                 270

tcc ctc agg aag att aat gct gcc tgg ttt aaa aat atg cct cat ctg   912
Ser Leu Arg Lys Ile Asn Ala Ala Trp Phe Lys Asn Met Pro His Leu
            275                 280                 285

aag gtg ctg gat ctt gaa ttc aac tat tta gtg gga gaa ata gcc tct   960
Lys Val Leu Asp Leu Glu Phe Asn Tyr Leu Val Gly Glu Ile Ala Ser
        290                 295                 300

ggg gca ttt tta acg atg ctg ccc cgc tta gaa ata ctt gac ttg tct  1008
Gly Ala Phe Leu Thr Met Leu Pro Arg Leu Glu Ile Leu Asp Leu Ser
    305                 310                 315

ttt aac tat ata aag ggg agt tat cca cag cat att aat att tcc aga  1056
Phe Asn Tyr Ile Lys Gly Ser Tyr Pro Gln His Ile Asn Ile Ser Arg
320                 325                 330                 335

aac ttc tct aaa ctt ttg tct cta cgg gca ttg cat tta aga ggt tat  1104
Asn Phe Ser Lys Leu Leu Ser Leu Arg Ala Leu His Leu Arg Gly Tyr
                340                 345                 350

gtg ttc cag gaa ctc aga gaa gat gat ttc cag ccc ctg atg cag ctt  1152
Val Phe Gln Glu Leu Arg Glu Asp Asp Phe Gln Pro Leu Met Gln Leu
            355                 360                 365

cca aac tta tcg act atc aac ttg ggt att aat ttt att aag caa atc  1200
Pro Asn Leu Ser Thr Ile Asn Leu Gly Ile Asn Phe Ile Lys Gln Ile
        370                 375                 380

gat ttc aaa ctt ttc caa aat ttc tcc aat ctg gaa att att tac ttg  1248
Asp Phe Lys Leu Phe Gln Asn Phe Ser Asn Leu Glu Ile Ile Tyr Leu
    385                 390                 395
```

```
tca gaa aac aga ata tca ccg ttg gta aaa gat acc cgg cag agt tat   1296
Ser Glu Asn Arg Ile Ser Pro Leu Val Lys Asp Thr Arg Gln Ser Tyr
400                 405                 410                 415

gca aat agt tcc tct ttt caa cgt cat atc cgg aaa cga cgc tca aca   1344
Ala Asn Ser Ser Ser Phe Gln Arg His Ile Arg Lys Arg Arg Ser Thr
                420                 425                 430

gat ttt gag ttt gac cca cat tcg aac ttt tat cat ttc acc cgt cct   1392
Asp Phe Glu Phe Asp Pro His Ser Asn Phe Tyr His Phe Thr Arg Pro
            435                 440                 445

tta ata aag cca caa tgt gct gct tat gga aaa gcc tta gat tta agc   1440
Leu Ile Lys Pro Gln Cys Ala Ala Tyr Gly Lys Ala Leu Asp Leu Ser
        450                 455                 460

ctc aac agt att ttc ttc att ggg cca aac caa ttt gaa aat ctt cct   1488
Leu Asn Ser Ile Phe Phe Ile Gly Pro Asn Gln Phe Glu Asn Leu Pro
    465                 470                 475

gac att gcc tgt tta aat ctg tct gca aat agc aat gct caa gtg tta   1536
Asp Ile Ala Cys Leu Asn Leu Ser Ala Asn Ser Asn Ala Gln Val Leu
480                 485                 490                 495

agt gga act gaa ttt tca gcc att cct cat gtc aaa tat ttg gat ttg   1584
Ser Gly Thr Glu Phe Ser Ala Ile Pro His Val Lys Tyr Leu Asp Leu
                500                 505                 510

aca aac aat aga cta gac ttt gat aat gct agt gct ctt act gaa ttg   1632
Thr Asn Asn Arg Leu Asp Phe Asp Asn Ala Ser Ala Leu Thr Glu Leu
            515                 520                 525

tcc gac ttg gaa gtt cta gat ctc agc tat aat tca cac tat ttc aga   1680
Ser Asp Leu Glu Val Leu Asp Leu Ser Tyr Asn Ser His Tyr Phe Arg
        530                 535                 540

ata gca ggc gta aca cat cat cta gaa ttt att caa aat ttc aca aat   1728
Ile Ala Gly Val Thr His His Leu Glu Phe Ile Gln Asn Phe Thr Asn
    545                 550                 555

cta aaa gtt tta aac ttg agc cac aac aac att tat act tta aca gat   1776
Leu Lys Val Leu Asn Leu Ser His Asn Asn Ile Tyr Thr Leu Thr Asp
560                 565                 570                 575

aag tat aac ctg gaa agc aag tcc ctg gta gaa tta gtt ttc agt ggc   1824
Lys Tyr Asn Leu Glu Ser Lys Ser Leu Val Glu Leu Val Phe Ser Gly
                580                 585                 590

aat cgc ctt gac att ttg tgg aat gat gat gac aac agg tat atc tcc   1872
Asn Arg Leu Asp Ile Leu Trp Asn Asp Asp Asp Asn Arg Tyr Ile Ser
            595                 600                 605

att ttc aaa ggt ctc aag aat ctg aca cgt ctg gat tta tcc ctt aat   1920
Ile Phe Lys Gly Leu Lys Asn Leu Thr Arg Leu Asp Leu Ser Leu Asn
        610                 615                 620

agg ctc aag cac atc cca aat gaa gca ttc ctt aat ttg cca gcg agt   1968
```

```
Arg Leu Lys His Ile Pro Asn Glu Ala Phe Leu Asn Leu Pro Ala Ser
625                 630                 635

ctc act gaa cta cat ata aat gat aat atg tta aag ttt ttt aac tgg   2016
Leu Thr Glu Leu His Ile Asn Asp Asn Met Leu Lys Phe Phe Asn Trp
640                 645                 650                 655

aca tta ctc cag cag ttt cct cgt ctc gag ttg ctt gac tta cgt gga   2064
Thr Leu Leu Gln Gln Phe Pro Arg Leu Glu Leu Leu Asp Leu Arg Gly
                660                 665                 670

aac aaa cta ctc ttt tta act gat agc cta tct gac ttt aca tct tcc   2112
Asn Lys Leu Leu Phe Leu Thr Asp Ser Leu Ser Asp Phe Thr Ser Ser
            675                 680                 685

ctt cgg aca ctg ctg ctg agt cat aac agg att tcc cac cta ccc tct   2160
Leu Arg Thr Leu Leu Leu Ser His Asn Arg Ile Ser His Leu Pro Ser
        690                 695                 700

ggc ttt ctt tct gaa gtc agt agt ctg aag cac ctc gat tta agt tcc   2208
Gly Phe Leu Ser Glu Val Ser Ser Leu Lys His Leu Asp Leu Ser Ser
705                 710                 715

aat ctg cta aaa aca atm aac aaa tcc gca ctt gaa act aag acc acc   2256
Asn Leu Leu Lys Thr Xaa Asn Lys Ser Ala Leu Glu Thr Lys Thr Thr
720                 725                 730                 735

acc aaa tta tct atg ttg gaa cta cac gga aac ccc ttt gaa tgc acc   2304
Thr Lys Leu Ser Met Leu Glu Leu His Gly Asn Pro Phe Glu Cys Thr
                740                 745                 750

tgt gac att gga gat ttc cga aga tgg atg gat gaa cat ctg aat gtc   2352
Cys Asp Ile Gly Asp Phe Arg Arg Trp Met Asp Glu His Leu Asn Val
            755                 760                 765

aaa att ccc aga ctg gta gat gtc att tgt gcc agt cct ggg gat caa   2400
Lys Ile Pro Arg Leu Val Asp Val Ile Cys Ala Ser Pro Gly Asp Gln
        770                 775                 780

aga ggg aag agt att gtg agt ctg gag cta aca act tgt gtt tca gat   2448
Arg Gly Lys Ser Ile Val Ser Leu Glu Leu Thr Thr Cys Val Ser Asp
785                 790                 795

gtc act gca gtg ata tta ttt ttc ttc acg ttc ttt atc acc acc atg   2496
Val Thr Ala Val Ile Leu Phe Phe Phe Thr Phe Phe Ile Thr Thr Met
800                 805                 810                 815

gtt atg ttg gct gcc ctg gct cac cat ttg ttt tac tgg gat gtt tgg   2544
Val Met Leu Ala Ala Leu Ala His His Leu Phe Tyr Trp Asp Val Trp
                820                 825                 830

ttt ata tat aat gtg tgt tta gct aag tta aaa ggc tac agg tct ctt   2592
Phe Ile Tyr Asn Val Cys Leu Ala Lys Leu Lys Gly Tyr Arg Ser Leu
            835                 840                 845

tcc aca tcc caa act ttc tat gat gct tac att tct tat gac acc aaa   2640
Ser Thr Ser Gln Thr Phe Tyr Asp Ala Tyr Ile Ser Tyr Asp Thr Lys
        850                 855                 860
```

```
gat gcc tct gtt act gac tgg gtg ata aat gag ctg cgc tac cac ctt   2688
Asp Ala Ser Val Thr Asp Trp Val Ile Asn Glu Leu Arg Tyr His Leu
    865                 870                 875

gaa gag agc cga gac aaa aac gtt ctc ctt tgt cta gag gag agg gat   2736
Glu Glu Ser Arg Asp Lys Asn Val Leu Leu Cys Leu Glu Glu Arg Asp
880                 885                 890                 895

tgg gac ccg gga ttg gcc atc atc gac aac ctc atg cag agc atc aac   2784
Trp Asp Pro Gly Leu Ala Ile Ile Asp Asn Leu Met Gln Ser Ile Asn
                900                 905                 910

caa agc aag aaa aca gta ttt gtt tta acc aaa aaa tat gca aaa agc   2832
Gln Ser Lys Lys Thr Val Phe Val Leu Thr Lys Lys Tyr Ala Lys Ser
            915                 920                 925

tgg aac ttt aaa aca gct ttt tac ttg gcc ttg cag agg cta atg ggt   2880
Trp Asn Phe Lys Thr Ala Phe Tyr Leu Ala Leu Gln Arg Leu Met Gly
        930                 935                 940

gag aac atg gat gtg att ata ttt atc ctg ctg gag cca gtg tta cag   2928
Glu Asn Met Asp Val Ile Ile Phe Ile Leu Leu Glu Pro Val Leu Gln
    945                 950                 955

cat tct ccg tat ttg agg cta cgg cag cgg atc tgt aag agc tcc atc   2976
His Ser Pro Tyr Leu Arg Leu Arg Gln Arg Ile Cys Lys Ser Ser Ile
960                 965                 970                 975

ctc cag tgg cct gac aac ccg aag gca gaa ggc ttg ttt tgg caa act   3024
Leu Gln Trp Pro Asp Asn Pro Lys Ala Glu Gly Leu Phe Trp Gln Thr
                980                 985                 990

ctg aga aat gtg gtc ttg act gaa aat gat tca cgg tat aac aat atg   3072
Leu Arg Asn Val Val Leu Thr Glu Asn Asp Ser Arg Tyr Asn Asn Met
            995                1000                1005

tat gtc gat tcc att aag caa tac taa                             3099
Tyr Val Asp Ser Ile Lys Gln Tyr
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Asn Asp Ser Val Ile Ala Glu Cys Ser Asn Arg Arg Leu Gln Glu Val
                20                  25                  30

Pro Gln Thr Val Gly Lys Tyr Val Thr Glu Leu Asp Leu Ser Asp Asn
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35 40 45

Phe Ile Thr His Ile Thr Asn Glu Ser Phe Gln Gly Leu Gln Asn Leu
50 55 60

Thr Lys Ile Asn Leu Asn His Asn Pro Asn Val Gln His Gln Asn Gly
65 70 75

Asn Pro Gly Ile Gln Ser Asn Gly Leu Asn Ile Thr Asp Gly Ala Phe
80 85 90 95

Leu Asn Leu Lys Asn Leu Arg Glu Leu Leu Leu Glu Asp Asn Gln Leu
100 105 110

Pro Gln Ile Pro Ser Gly Leu Pro Glu Ser Leu Thr Glu Leu Ser Leu
115 120 125

Ile Gln Asn Asn Ile Tyr Asn Ile Thr Lys Glu Gly Ile Ser Arg Leu
130 135 140

Ile Asn Leu Lys Asn Leu Tyr Leu Ala Trp Asn Cys Tyr Phe Asn Lys
145 150 155

Val Cys Glu Lys Thr Asn Ile Glu Asp Gly Val Phe Glu Thr Leu Thr
160 165 170 175

Asn Leu Glu Leu Leu Ser Leu Ser Phe Asn Ser Leu Ser His Val Pro
180 185 190

Pro Lys Leu Pro Ser Ser Leu Arg Lys Leu Phe Leu Ser Asn Thr Gln
195 200 205

Ile Lys Tyr Ile Ser Glu Glu Asp Phe Lys Gly Leu Ile Asn Leu Thr
210 215 220

Leu Leu Asp Leu Ser Gly Asn Cys Pro Arg Cys Phe Asn Ala Pro Phe
225 230 235

Pro Cys Val Pro Cys Asp Gly Gly Ala Ser Ile Asn Ile Asp Arg Phe
240 245 250 255

Ala Phe Gln Asn Leu Thr Gln Leu Arg Tyr Leu Asn Leu Ser Ser Thr
260 265 270

Ser Leu Arg Lys Ile Asn Ala Ala Trp Phe Lys Asn Met Pro His Leu
275 280 285

Lys Val Leu Asp Leu Glu Phe Asn Tyr Leu Val Gly Glu Ile Ala Ser
290 295 300

Gly Ala Phe Leu Thr Met Leu Pro Arg Leu Glu Ile Leu Asp Leu Ser
305 310 315

Phe Asn Tyr Ile Lys Gly Ser Tyr Pro Gln His Ile Asn Ile Ser Arg
320 325 330 335

Asn Phe Ser Lys Leu Leu Ser Leu Arg Ala Leu His Leu Arg Gly Tyr
340 345 350

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Val Phe Gln Glu Leu Arg Glu Asp Asp Phe Gln Pro Leu Met Gln Leu
            355                 360                 365

Pro Asn Leu Ser Thr Ile Asn Leu Gly Ile Asn Phe Ile Lys Gln Ile
        370                 375                 380

Asp Phe Lys Leu Phe Gln Asn Phe Ser Asn Leu Glu Ile Ile Tyr Leu
    385                 390                 395

Ser Glu Asn Arg Ile Ser Pro Leu Val Lys Asp Thr Arg Gln Ser Tyr
400                 405                 410                 415

Ala Asn Ser Ser Ser Phe Gln Arg His Ile Arg Lys Arg Arg Ser Thr
                420                 425                 430

Asp Phe Glu Phe Asp Pro His Ser Asn Phe Tyr His Phe Thr Arg Pro
            435                 440                 445

Leu Ile Lys Pro Gln Cys Ala Ala Tyr Gly Lys Ala Leu Asp Leu Ser
        450                 455                 460

Leu Asn Ser Ile Phe Phe Ile Gly Pro Asn Gln Phe Glu Asn Leu Pro
    465                 470                 475

Asp Ile Ala Cys Leu Asn Leu Ser Ala Asn Ser Asn Ala Gln Val Leu
480                 485                 490                 495

Ser Gly Thr Glu Phe Ser Ala Ile Pro His Val Lys Tyr Leu Asp Leu
                500                 505                 510

Thr Asn Asn Arg Leu Asp Phe Asp Asn Ala Ser Ala Leu Thr Glu Leu
            515                 520                 525

Ser Asp Leu Glu Val Leu Asp Leu Ser Tyr Asn Ser His Tyr Phe Arg
        530                 535                 540

Ile Ala Gly Val Thr His His Leu Glu Phe Ile Gln Asn Phe Thr Asn
    545                 550                 555

Leu Lys Val Leu Asn Leu Ser His Asn Asn Ile Tyr Thr Leu Thr Asp
560                 565                 570                 575

Lys Tyr Asn Leu Glu Ser Lys Ser Leu Val Glu Leu Val Phe Ser Gly
                580                 585                 590

Asn Arg Leu Asp Ile Leu Trp Asn Asp Asp Asp Asn Arg Tyr Ile Ser
            595                 600                 605

Ile Phe Lys Gly Leu Lys Asn Leu Thr Arg Leu Asp Leu Ser Leu Asn
        610                 615                 620

Arg Leu Lys His Ile Pro Asn Glu Ala Phe Leu Asn Leu Pro Ala Ser
    625                 630                 635

Leu Thr Glu Leu His Ile Asn Asp Asn Met Leu Lys Phe Phe Asn Trp
640                 645                 650                 655
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Thr Leu Leu Gln Gln Phe Pro Arg Leu Glu Leu Leu Asp Leu Arg Gly
660 665 670

Asn Lys Leu Leu Phe Leu Thr Asp Ser Leu Ser Asp Phe Thr Ser Ser
675 680 685

Leu Arg Thr Leu Leu Leu Ser His Asn Arg Ile Ser His Leu Pro Ser
690 695 700

Gly Phe Leu Ser Glu Val Ser Ser Leu Lys His Leu Asp Leu Ser Ser
705 710 715

Asn Leu Leu Lys Thr Xaa Asn Lys Ser Ala Leu Glu Thr Lys Thr Thr
720 725 730 735

Thr Lys Leu Ser Met Leu Glu Leu His Gly Asn Pro Phe Glu Cys Thr
740 745 750

Cys Asp Ile Gly Asp Phe Arg Arg Trp Met Asp Glu His Leu Asn Val
755 760 765

Lys Ile Pro Arg Leu Val Asp Val Ile Cys Ala Ser Pro Gly Asp Gln
770 775 780

Arg Gly Lys Ser Ile Val Ser Leu Glu Leu Thr Thr Cys Val Ser Asp
785 790 795

Val Thr Ala Val Ile Leu Phe Phe Phe Thr Phe Phe Ile Thr Thr Met
800 805 810 815

Val Met Leu Ala Ala Leu Ala His His Leu Phe Tyr Trp Asp Val Trp
820 825 830

Phe Ile Tyr Asn Val Cys Leu Ala Lys Leu Lys Gly Tyr Arg Ser Leu
835 840 845

Ser Thr Ser Gln Thr Phe Tyr Asp Ala Tyr Ile Ser Tyr Asp Thr Lys
850 855 860

Asp Ala Ser Val Thr Asp Trp Val Ile Asn Glu Leu Arg Tyr His Leu
865 870 875

Glu Glu Ser Arg Asp Lys Asn Val Leu Leu Cys Leu Glu Glu Arg Asp
880 885 890 895

Trp Asp Pro Gly Leu Ala Ile Ile Asp Asn Leu Met Gln Ser Ile Asn
900 905 910

Gln Ser Lys Lys Thr Val Phe Val Leu Thr Lys Lys Tyr Ala Lys Ser
915 920 925

Trp Asn Phe Lys Thr Ala Phe Tyr Leu Ala Leu Gln Arg Leu Met Gly
930 935 940

Glu Asn Met Asp Val Ile Ile Phe Ile Leu Leu Glu Pro Val Leu Gln
945 950 955

His Ser Pro Tyr Leu Arg Leu Arg Gln Arg Ile Cys Lys Ser Ser Ile

960 965 970 975

Leu Gln Trp Pro Asp Asn Pro Lys Ala Glu Gly Leu Phe Trp Gln Thr
980 985 990

Leu Arg Asn Val Val Leu Thr Glu Asn Asp Ser Arg Tyr Asn Asn Met
995 1000 1005

Tyr Val Asp Ser Ile Lys Gln Tyr
1010 1015

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<222> (168)..(2543)

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atcttcatca ttcatatgag gaaataagtg gtaaaatcct tggaaataca atg aga 116
Met Arg

ctc atc aga aac att tac ata ttt tgt agt att gtt atg aca gca gag 164
Leu Ile Arg Asn Ile Tyr Ile Phe Cys Ser Ile Val Met Thr Ala Glu
-15 -10 -5

ggt gat gct cca gag ctg cca gaa gaa agg gaa ctg atg acc aac tgc 212
Gly Asp Ala Pro Glu Leu Pro Glu Glu Arg Glu Leu Met Thr Asn Cys
-1 1 5 10 15

tcc aac atg tct cta aga aag gtt ccc gca gac ttg acc cca gcc aca 260
Ser Asn Met Ser Leu Arg Lys Val Pro Ala Asp Leu Thr Pro Ala Thr
20 25 30

acg aca ctg gat tta tcc tat aac ctc ctt ttt caa ctc cag agt tca 308
Thr Thr Leu Asp Leu Ser Tyr Asn Leu Leu Phe Gln Leu Gln Ser Ser
35 40 45

gat ttt cat tct gtc tcc aaa ctg aga gtt ttg att cta tgc cat aac 356
Asp Phe His Ser Val Ser Lys Leu Arg Val Leu Ile Leu Cys His Asn
50 55 60

aga att caa cag ctg gat ctc aaa acc ttt gaa ttc aac aag gag tta 404
Arg Ile Gln Gln Leu Asp Leu Lys Thr Phe Glu Phe Asn Lys Glu Leu
65 70 75

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aga tat tta gat ttg tct aat aac aga ctg aag agt gta act tgg tat   452
Arg Tyr Leu Asp Leu Ser Asn Asn Arg Leu Lys Ser Val Thr Trp Tyr
 80                  85                  90                  95

tta ctg gca ggt ctc agg tat tta gat ctt tct ttt aat gac ttt gac   500
Leu Leu Ala Gly Leu Arg Tyr Leu Asp Leu Ser Phe Asn Asp Phe Asp
                100                 105                 110

acc atg cct atc tgt gag gaa gct ggc aac atg tca cac ctg gaa atc   548
Thr Met Pro Ile Cys Glu Glu Ala Gly Asn Met Ser His Leu Glu Ile
            115                 120                 125

cta ggt ttg agt ggg gca aaa ata caa aaa tca gat ttc cag aaa att   596
Leu Gly Leu Ser Gly Ala Lys Ile Gln Lys Ser Asp Phe Gln Lys Ile
        130                 135                 140

gct cat ctg cat cta aat act gtc ttc tta gga ttc aga act ctt cct   644
Ala His Leu His Leu Asn Thr Val Phe Leu Gly Phe Arg Thr Leu Pro
    145                 150                 155

cat tat gaa gaa ggt agc ctg ccc atc tta aac aca aca aaa ctg cac   692
His Tyr Glu Glu Gly Ser Leu Pro Ile Leu Asn Thr Thr Lys Leu His
160                 165                 170                 175

att gtt tta cca atg gac aca aat ttc tgg gtt ctt ttg cgt gat gga   740
Ile Val Leu Pro Met Asp Thr Asn Phe Trp Val Leu Leu Arg Asp Gly
                180                 185                 190

atc aag act tca aaa ata tta gaa atg aca aat ata gat ggc aaa agc   788
Ile Lys Thr Ser Lys Ile Leu Glu Met Thr Asn Ile Asp Gly Lys Ser
            195                 200                 205

caa ttt gta agt tat gaa atg caa cga aat ctt agt tta gaa aat gct   836
Gln Phe Val Ser Tyr Glu Met Gln Arg Asn Leu Ser Leu Glu Asn Ala
        210                 215                 220

aag aca tcg gtt cta ttg ctt aat aaa gtt gat tta ctc tgg gac gac   884
Lys Thr Ser Val Leu Leu Leu Asn Lys Val Asp Leu Leu Trp Asp Asp
    225                 230                 235

ctt ttc ctt atc tta caa ttt gtt tgg cat aca tca gtg gaa cac ttt   932
Leu Phe Leu Ile Leu Gln Phe Val Trp His Thr Ser Val Glu His Phe
240                 245                 250                 255

cag atc cga aat gtg act ttt ggt ggt aag gct tat ctt gac cac aat   980
Gln Ile Arg Asn Val Thr Phe Gly Gly Lys Ala Tyr Leu Asp His Asn
                260                 265                 270

tca ttt gac tac tca aat act gta atg aga act ata aaa ttg gag cat  1028
Ser Phe Asp Tyr Ser Asn Thr Val Met Arg Thr Ile Lys Leu Glu His
            275                 280                 285

gta cat ttc aga gtg ttt tac att caa cag gat aaa atc tat ttg ctt  1076
Val His Phe Arg Val Phe Tyr Ile Gln Gln Asp Lys Ile Tyr Leu Leu
        290                 295                 300

ttg acc aaa atg gac ata gaa aac ctg aca ata tca aat gca caa atg  1124
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Leu Thr Lys Met Asp Ile Glu Asn Leu Thr Ile Ser Asn Ala Gln Met
    305                 310                 315

cca cac atg ctt ttc ccg aat tat cct acg aaa ttc caa tat tta aat  1172
Pro His Met Leu Phe Pro Asn Tyr Pro Thr Lys Phe Gln Tyr Leu Asn
320                 325                 330                 335

ttt gcc aat aat atc tta aca gac gag ttg ttt aaa aga act atc caa  1220
Phe Ala Asn Asn Ile Leu Thr Asp Glu Leu Phe Lys Arg Thr Ile Gln
                340                 345                 350

ctg cct cac ttg aaa act ctc att ttg aat ggc aat aaa ctg gag aca  1268
Leu Pro His Leu Lys Thr Leu Ile Leu Asn Gly Asn Lys Leu Glu Thr
            355                 360                 365

ctt tct tta gta agt tgc ttt gct aac aac aca ccc ttg gaa cac ttg  1316
Leu Ser Leu Val Ser Cys Phe Ala Asn Asn Thr Pro Leu Glu His Leu
        370                 375                 380

gat ctg agt caa aat cta tta caa cat aaa aat gat gaa aat tgc tca  1364
Asp Leu Ser Gln Asn Leu Leu Gln His Lys Asn Asp Glu Asn Cys Ser
    385                 390                 395

tgg cca gaa act gtg gtc aat atg aat ctg tca tac aat aaa ttg tct  1412
Trp Pro Glu Thr Val Val Asn Met Asn Leu Ser Tyr Asn Lys Leu Ser
400                 405                 410                 415

gat tct gtc ttc agg tgc ttg ccc aaa agt att caa ata ctt gac cta  1460
Asp Ser Val Phe Arg Cys Leu Pro Lys Ser Ile Gln Ile Leu Asp Leu
                420                 425                 430

aat aat aac caa atc caa act gta cct aaa gag act att cat ctg atg  1508
Asn Asn Asn Gln Ile Gln Thr Val Pro Lys Glu Thr Ile His Leu Met
            435                 440                 445

gcc tta cga gaa cta aat att gca ttt aat ttt cta act gat ctc cct  1556
Ala Leu Arg Glu Leu Asn Ile Ala Phe Asn Phe Leu Thr Asp Leu Pro
        450                 455                 460

gga tgc agt cat ttc agt aga ctt tca gtt ctg aac att gaa atg aac  1604
Gly Cys Ser His Phe Ser Arg Leu Ser Val Leu Asn Ile Glu Met Asn
    465                 470                 475

ttc att ctc agc cca tct ctg gat ttt gtt cag agc tgc cag gaa gtt  1652
Phe Ile Leu Ser Pro Ser Leu Asp Phe Val Gln Ser Cys Gln Glu Val
480                 485                 490                 495

aaa act cta aat gcg gga aga aat cca ttc cgg tgt acc tgt gaa tta  1700
Lys Thr Leu Asn Ala Gly Arg Asn Pro Phe Arg Cys Thr Cys Glu Leu
                500                 505                 510

aaa aat ttc att cag ctt gaa aca tat tca gag gtc atg atg gtt gga  1748
Lys Asn Phe Ile Gln Leu Glu Thr Tyr Ser Glu Val Met Met Val Gly
            515                 520                 525

tgg tca gat tca tac acc tgt gaa tac cct tta aac cta agg gga act  1796
Trp Ser Asp Ser Tyr Thr Cys Glu Tyr Pro Leu Asn Leu Arg Gly Thr
        530                 535                 540
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agg tta aaa gac gtt cat ctc cac gaa tta tct tgc aac aca gct ctg   1844
Arg Leu Lys Asp Val His Leu His Glu Leu Ser Cys Asn Thr Ala Leu
    545                 550                 555

ttg att gtc acc att gtg gtt att atg cta gtt ctg ggg ttg gct gtg   1892
Leu Ile Val Thr Ile Val Val Ile Met Leu Val Leu Gly Leu Ala Val
560                 565                 570                 575

gcc ttc tgc tgt ctc cac ttt gat ctg ccc tgg tat ctc agg atg cta   1940
Ala Phe Cys Cys Leu His Phe Asp Leu Pro Trp Tyr Leu Arg Met Leu
                580                 585                 590

ggt caa tgc aca caa aca tgg cac agg gtt agg aaa aca acc caa gaa   1988
Gly Gln Cys Thr Gln Thr Trp His Arg Val Arg Lys Thr Thr Gln Glu
            595                 600                 605

caa ctc aag aga aat gtc cga ttc cac gca ttt att tca tac agt gaa   2036
Gln Leu Lys Arg Asn Val Arg Phe His Ala Phe Ile Ser Tyr Ser Glu
        610                 615                 620

cat gat tct ctg tgg gtg aag aat gaa ttg atc ccc aat cta gag aag   2084
His Asp Ser Leu Trp Val Lys Asn Glu Leu Ile Pro Asn Leu Glu Lys
    625                 630                 635

gaa gat ggt tct atc ttg att tgc ctt tat gaa agc tac ttt gac cct   2132
Glu Asp Gly Ser Ile Leu Ile Cys Leu Tyr Glu Ser Tyr Phe Asp Pro
640                 645                 650                 655

ggc aaa agc att agt gaa aat att gta agc ttc att gag aaa agc tat   2180
Gly Lys Ser Ile Ser Glu Asn Ile Val Ser Phe Ile Glu Lys Ser Tyr
                660                 665                 670

aag tcc atc ttt gtt ttg tct ccc aac ttt gtc cag aat gag tgg tgc   2228
Lys Ser Ile Phe Val Leu Ser Pro Asn Phe Val Gln Asn Glu Trp Cys
            675                 680                 685

cat tat gaa ttc tac ttt gcc cac cac aat ctc ttc cat gaa aat tct   2276
His Tyr Glu Phe Tyr Phe Ala His His Asn Leu Phe His Glu Asn Ser
        690                 695                 700

gat cat ata att ctt atc tta ctg gaa ccc att cca ttc tat tgc att   2324
Asp His Ile Ile Leu Ile Leu Leu Glu Pro Ile Pro Phe Tyr Cys Ile
    705                 710                 715

ccc acc agg tat cat aaa ctg aaa gct ctc ctg gaa aaa aaa gca tac   2372
Pro Thr Arg Tyr His Lys Leu Lys Ala Leu Leu Glu Lys Lys Ala Tyr
720                 725                 730                 735

ttg gaa tgg ccc aag gat agg cgt aaa tgt ggg ctt ttc tgg gca aac   2420
Leu Glu Trp Pro Lys Asp Arg Arg Lys Cys Gly Leu Phe Trp Ala Asn
                740                 745                 750

ctt cga gct gct att aat gtt aat gta tta gcc acc aga gaa atg tat   2468
Leu Arg Ala Ala Ile Asn Val Asn Val Leu Ala Thr Arg Glu Met Tyr
            755                 760                 765

gaa ctg cag aca ttc aca gag tta aat gaa gag tct cga ggt tct aca   2516
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Glu Leu Gln Thr Phe Thr Glu Leu Asn Glu Glu Ser Arg Gly Ser Thr
770 775 780

atc tct ctg atg aga aca gat tgt cta taaaatccca cagtccttgg 2563
Ile Ser Leu Met Arg Thr Asp Cys Leu
785 790

gaagttgggg accacataca ctgttgggat gtacattgat acaaccttta tgatggcaat 2623

ttgacaatat ttattaaaat aaaaaatggt tattcccttc atatcagttt ctagaaggat 2683

ttctaagaat gtatcctata gaaacacctt cacaagttta taagggctta tggaaaaagg 2743

tgttcatccc aggattgttt ataatcatga aaaatgtggc caggtgcagt ggctcactct 2803

tgtaatccca gcactatggg aggccaaggt gggtgaccca cgaggtcaag agatggagac 2863

catcctggcc aacatggtga aaccctgtct ctactaaaaa tacaaaaatt agctgggcgt 2923

gatggtgcac gcctgtagtc ccagctactt gggaggctga ggcaggagaa tcgcttgaac 2983

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Asn Cys Ser Asn Met Ser Leu Arg Lys Val Pro Ala Asp Leu Thr Pro
15 20 25

Ala Thr Thr Thr Leu Asp Leu Ser Tyr Asn Leu Leu Phe Gln Leu Gln
30 35 40 45

Ser Ser Asp Phe His Ser Val Ser Lys Leu Arg Val Leu Ile Leu Cys
50 55 60

His Asn Arg Ile Gln Gln Leu Asp Leu Lys Thr Phe Glu Phe Asn Lys
65 70 75

Glu Leu Arg Tyr Leu Asp Leu Ser Asn Asn Arg Leu Lys Ser Val Thr
80 85 90

Trp Tyr Leu Leu Ala Gly Leu Arg Tyr Leu Asp Leu Ser Phe Asn Asp
95 100 105

Phe Asp Thr Met Pro Ile Cys Glu Glu Ala Gly Asn Met Ser His Leu
110 115 120 125

Glu Ile Leu Gly Leu Ser Gly Ala Lys Ile Gln Lys Ser Asp Phe Gln
130 135 140

Lys Ile Ala His Leu His Leu Asn Thr Val Phe Leu Gly Phe Arg Thr
145 150 155

Leu Pro His Tyr Glu Glu Gly Ser Leu Pro Ile Leu Asn Thr Thr Lys
160 165 170

Leu His Ile Val Leu Pro Met Asp Thr Asn Phe Trp Val Leu Leu Arg
175 180 185

Asp Gly Ile Lys Thr Ser Lys Ile Leu Glu Met Thr Asn Ile Asp Gly
190 195 200 205

Lys Ser Gln Phe Val Ser Tyr Glu Met Gln Arg Asn Leu Ser Leu Glu
210 215 220

Asn Ala Lys Thr Ser Val Leu Leu Leu Asn Lys Val Asp Leu Leu Trp
225 230 235

Asp Asp Leu Phe Leu Ile Leu Gln Phe Val Trp His Thr Ser Val Glu
240 245 250

His Phe Gln Ile Arg Asn Val Thr Phe Gly Gly Lys Ala Tyr Leu Asp
255 260 265

His Asn Ser Phe Asp Tyr Ser Asn Thr Val Met Arg Thr Ile Lys Leu
270 275 280 285

Glu His Val His Phe Arg Val Phe Tyr Ile Gln Gln Asp Lys Ile Tyr
290 295 300

Leu Leu Leu Thr Lys Met Asp Ile Glu Asn Leu Thr Ile Ser Asn Ala
305 310 315

Gln Met Pro His Met Leu Phe Pro Asn Tyr Pro Thr Lys Phe Gln Tyr
320 325 330

Leu Asn Phe Ala Asn Asn Ile Leu Thr Asp Glu Leu Phe Lys Arg Thr
335 340 345

Ile Gln Leu Pro His Leu Lys Thr Leu Ile Leu Asn Gly Asn Lys Leu
350 355 360 365

Glu Thr Leu Ser Leu Val Ser Cys Phe Ala Asn Asn Thr Pro Leu Glu
370 375 380

His Leu Asp Leu Ser Gln Asn Leu Leu Gln His Lys Asn Asp Glu Asn
385 390 395

Cys Ser Trp Pro Glu Thr Val Val Asn Met Asn Leu Ser Tyr Asn Lys
400 405 410

Leu Ser Asp Ser Val Phe Arg Cys Leu Pro Lys Ser Ile Gln Ile Leu
415 420 425

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Asp Leu Asn Asn Asn Gln Ile Gln Thr Val Pro Lys Glu Thr Ile His
430                 435                 440                 445

Leu Met Ala Leu Arg Glu Leu Asn Ile Ala Phe Asn Phe Leu Thr Asp
                450                 455                 460

Leu Pro Gly Cys Ser His Phe Ser Arg Leu Ser Val Leu Asn Ile Glu
            465                 470                 475

Met Asn Phe Ile Leu Ser Pro Ser Leu Asp Phe Val Gln Ser Cys Gln
        480                 485                 490

Glu Val Lys Thr Leu Asn Ala Gly Arg Asn Pro Phe Arg Cys Thr Cys
    495                 500                 505

Glu Leu Lys Asn Phe Ile Gln Leu Glu Thr Tyr Ser Glu Val Met Met
510                 515                 520                 525

Val Gly Trp Ser Asp Ser Tyr Thr Cys Glu Tyr Pro Leu Asn Leu Arg
                530                 535                 540

Gly Thr Arg Leu Lys Asp Val His Leu His Glu Leu Ser Cys Asn Thr
            545                 550                 555

Ala Leu Leu Ile Val Thr Ile Val Val Ile Met Leu Val Leu Gly Leu
        560                 565                 570

Ala Val Ala Phe Cys Cys Leu His Phe Asp Leu Pro Trp Tyr Leu Arg
    575                 580                 585

Met Leu Gly Gln Cys Thr Gln Thr Trp His Arg Val Arg Lys Thr Thr
590                 595                 600                 605

Gln Glu Gln Leu Lys Arg Asn Val Arg Phe His Ala Phe Ile Ser Tyr
                610                 615                 620

Ser Glu His Asp Ser Leu Trp Val Lys Asn Glu Leu Ile Pro Asn Leu
            625                 630                 635

Glu Lys Glu Asp Gly Ser Ile Leu Ile Cys Leu Tyr Glu Ser Tyr Phe
        640                 645                 650

Asp Pro Gly Lys Ser Ile Ser Glu Asn Ile Val Ser Phe Ile Glu Lys
    655                 660                 665

Ser Tyr Lys Ser Ile Phe Val Leu Ser Pro Asn Phe Val Gln Asn Glu
670                 675                 680                 685

Trp Cys His Tyr Glu Phe Tyr Phe Ala His His Asn Leu Phe His Glu
                690                 695                 700

Asn Ser Asp His Ile Ile Leu Ile Leu Leu Glu Pro Ile Pro Phe Tyr
            705                 710                 715

Cys Ile Pro Thr Arg Tyr His Lys Leu Lys Ala Leu Leu Glu Lys Lys
        720                 725                 730

Ala Tyr Leu Glu Trp Pro Lys Asp Arg Arg Lys Cys Gly Leu Phe Trp
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735 740 745

Ala Asn Leu Arg Ala Ala Ile Asn Val Asn Val Leu Ala Thr Arg Glu
750 755 760 765

Met Tyr Glu Leu Gln Thr Phe Thr Glu Leu Asn Glu Glu Ser Arg Gly
770 775 780

Ser Thr Ile Ser Leu Met Arg Thr Asp Cys Leu
785 790

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Met Thr Lys Asp Lys Glu Pro Ile Val Lys Ser Phe His Phe
-30 -25 -20

gtt tgc ctt atg atc ata ata gtt gga acc aga atc cag ttc tcc gac 157
Val Cys Leu Met Ile Ile Ile Val Gly Thr Arg Ile Gln Phe Ser Asp
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gga aat gaa ttt gca gta gac aag tca aaa aga ggt ctt att cat gtt 205
Gly Asn Glu Phe Ala Val Asp Lys Ser Lys Arg Gly Leu Ile His Val
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cca aaa gac cta ccg ctg aaa acc aaa gtc tta gat atg tct cag aac 253
Pro Lys Asp Leu Pro Leu Lys Thr Lys Val Leu Asp Met Ser Gln Asn
20 25 30

tac atc gct gag ctt cag gtc tct gac atg agc ttt cta tca gag ttg 301
Tyr Ile Ala Glu Leu Gln Val Ser Asp Met Ser Phe Leu Ser Glu Leu
35 40 45

aca gtt ttg aga ctt tcc cat aac aga atc cag cta ctt gat tta agt 349

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Thr Val Leu Arg Leu Ser His Asn Arg Ile Gln Leu Leu Asp Leu Ser
          50                  55                  60

gtt ttc aag ttc aac cag gat tta gaa tat ttg gat tta tct cat aat   397
Val Phe Lys Phe Asn Gln Asp Leu Glu Tyr Leu Asp Leu Ser His Asn
 65                  70                  75

cag ttg caa aag ata tcc tgc cat cct att gtg agt ttc agg cat tta   445
Gln Leu Gln Lys Ile Ser Cys His Pro Ile Val Ser Phe Arg His Leu
 80                  85                  90                  95

gat ctc tca ttc aat gat ttc aag gcc ctg ccc atc tgt aag gaa ttt   493
Asp Leu Ser Phe Asn Asp Phe Lys Ala Leu Pro Ile Cys Lys Glu Phe
                100                 105                 110

ggc aac tta tca caa ctg aat ttc ttg gga ttg agt gct atg aag ctg   541
Gly Asn Leu Ser Gln Leu Asn Phe Leu Gly Leu Ser Ala Met Lys Leu
            115                 120                 125

caa aaa tta gat ttg ctg cca att gct cac ttg cat cta agt tat atc   589
Gln Lys Leu Asp Leu Leu Pro Ile Ala His Leu His Leu Ser Tyr Ile
        130                 135                 140

ctt ctg gat tta aga aat tat tat ata aaa gaa aat gag aca gaa agt   637
Leu Leu Asp Leu Arg Asn Tyr Tyr Ile Lys Glu Asn Glu Thr Glu Ser
    145                 150                 155

cta caa att ctg aat gca aaa acc ctt cac ctt gtt ttt cac cca act   685
Leu Gln Ile Leu Asn Ala Lys Thr Leu His Leu Val Phe His Pro Thr
160                 165                 170                 175

agt tta ttc gct atc caa gtg aac ata tca gtt aat act tta ggg tgc   733
Ser Leu Phe Ala Ile Gln Val Asn Ile Ser Val Asn Thr Leu Gly Cys
                180                 185                 190

tta caa ctg act aat att aaa ttg aat gat gac aac tgt caa gtt ttc   781
Leu Gln Leu Thr Asn Ile Lys Leu Asn Asp Asp Asn Cys Gln Val Phe
            195                 200                 205

att aaa ttt tta tca gaa ctc acc aga ggt cca acc tta ctg aat ttt   829
Ile Lys Phe Leu Ser Glu Leu Thr Arg Gly Pro Thr Leu Leu Asn Phe
        210                 215                 220

acc ctc aac cac ata gaa acg act tgg aaa tgc ctg gtc aga gtc ttt   877
Thr Leu Asn His Ile Glu Thr Thr Trp Lys Cys Leu Val Arg Val Phe
    225                 230                 235

caa ttt ctt tgg ccc aaa cct gtg gaa tat ctc aat att tac aat tta   925
Gln Phe Leu Trp Pro Lys Pro Val Glu Tyr Leu Asn Ile Tyr Asn Leu
240                 245                 250                 255

aca ata att gaa agc att cgt gaa gaa gat ttt act tat tct aaa acg   973
Thr Ile Ile Glu Ser Ile Arg Glu Glu Asp Phe Thr Tyr Ser Lys Thr
                260                 265                 270

aca ttg aaa gca ttg aca ata gaa cat atc acg aac caa gtt ttt ctg  1021
Thr Leu Lys Ala Leu Thr Ile Glu His Ile Thr Asn Gln Val Phe Leu
            275                 280                 285
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ttt tca cag aca gct ttg tac acc gtg ttt tct gag atg aac att atg   1069
Phe Ser Gln Thr Ala Leu Tyr Thr Val Phe Ser Glu Met Asn Ile Met
        290                 295                 300

atg tta acc att tca gat aca cct ttt ata cac atg ctg tgt cct cat   1117
Met Leu Thr Ile Ser Asp Thr Pro Phe Ile His Met Leu Cys Pro His
    305                 310                 315

gca cca agc aca ttc aag ttt ttg aac ttt acc cag aac gtt ttc aca   1165
Ala Pro Ser Thr Phe Lys Phe Leu Asn Phe Thr Gln Asn Val Phe Thr
320                 325                 330                 335

gat agt att ttt gaa aaa tgt tcc acg tta gtt aaa ttg gag aca ctt   1213
Asp Ser Ile Phe Glu Lys Cys Ser Thr Leu Val Lys Leu Glu Thr Leu
                340                 345                 350

atc tta caa aag aat gga tta aaa gac ctt ttc aaa gta ggt ctc atg   1261
Ile Leu Gln Lys Asn Gly Leu Lys Asp Leu Phe Lys Val Gly Leu Met
            355                 360                 365

acg aag gat atg cct tct ttg gaa ata ctg gat gtt agc tgg aat tct   1309
Thr Lys Asp Met Pro Ser Leu Glu Ile Leu Asp Val Ser Trp Asn Ser
        370                 375                 380

ttg gaa tct ggt aga cat aaa gaa aac tgc act tgg gtt gag agt ata   1357
Leu Glu Ser Gly Arg His Lys Glu Asn Cys Thr Trp Val Glu Ser Ile
    385                 390                 395

gtg gtg tta aat ttg tct tca aat atg ctt act gac tct gtt ttc aga   1405
Val Val Leu Asn Leu Ser Ser Asn Met Leu Thr Asp Ser Val Phe Arg
400                 405                 410                 415

tgt tta cct ccc agg atc aag gta ctt gat ctt cac agc aat aaa ata   1453
Cys Leu Pro Pro Arg Ile Lys Val Leu Asp Leu His Ser Asn Lys Ile
                420                 425                 430

aag agc gtt cct aaa caa gtc gta aaa ctg gaa gct ttg caa gaa ctc   1501
Lys Ser Val Pro Lys Gln Val Val Lys Leu Glu Ala Leu Gln Glu Leu
            435                 440                 445

aat gtt gct ttc aat tct tta act gac ctt cct gga tgt ggc agc ttt   1549
Asn Val Ala Phe Asn Ser Leu Thr Asp Leu Pro Gly Cys Gly Ser Phe
        450                 455                 460

agc agc ctt tct gta ttg atc att gat cac aat tca gtt tcc cac cca   1597
Ser Ser Leu Ser Val Leu Ile Ile Asp His Asn Ser Val Ser His Pro
    465                 470                 475

tcg gct gat ttc ttc cag agc tgc cag aag atg agg tca ata aaa gca   1645
Ser Ala Asp Phe Phe Gln Ser Cys Gln Lys Met Arg Ser Ile Lys Ala
480                 485                 490                 495

ggg gac aat cca ttc caa tgt acc tgt gag cta aga gaa ttt gtc aaa   1693
Gly Asp Asn Pro Phe Gln Cys Thr Cys Glu Leu Arg Glu Phe Val Lys
                500                 505                 510

aat ata gac caa gta tca agt gaa gtg tta gag ggc tgg cct gat tct   1741
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Asn Ile Asp Gln Val Ser Ser Glu Val Leu Glu Gly Trp Pro Asp Ser
                515                 520                 525

tat aag tgt gac tac cca gaa agt tat aga gga agc cca cta aag gac  1789
Tyr Lys Cys Asp Tyr Pro Glu Ser Tyr Arg Gly Ser Pro Leu Lys Asp
        530                 535                 540

ttt cac atg tct gaa tta tcc tgc aac ata act ctg ctg atc gtc acc  1837
Phe His Met Ser Glu Leu Ser Cys Asn Ile Thr Leu Leu Ile Val Thr
    545                 550                 555

atc ggt gcc acc atg ctg gtg ttg gct gtg act gtg acc tcc ctc tgc  1885
Ile Gly Ala Thr Met Leu Val Leu Ala Val Thr Val Thr Ser Leu Cys
560                 565                 570                 575

atc tac ttg gat ctg ccc tgg tat ctc agg atg gtg tgc cag tgg acc  1933
Ile Tyr Leu Asp Leu Pro Trp Tyr Leu Arg Met Val Cys Gln Trp Thr
                580                 585                 590

cag act cgg cgc agg gcc agg aac ata ccc tta gaa gaa ctc caa aga  1981
Gln Thr Arg Arg Arg Ala Arg Asn Ile Pro Leu Glu Glu Leu Gln Arg
            595                 600                 605

aac ctc cag ttt cat gct ttt att tca tat agt gaa cat gat tct gcc  2029
Asn Leu Gln Phe His Ala Phe Ile Ser Tyr Ser Glu His Asp Ser Ala
        610                 615                 620

tgg gtg aaa agt gaa ttg gta cct tac cta gaa aaa gaa gat ata cag  2077
Trp Val Lys Ser Glu Leu Val Pro Tyr Leu Glu Lys Glu Asp Ile Gln
    625                 630                 635

att tgt ctt cat gag agg aac ttt gtc cct ggc aag agc att gtg gaa  2125
Ile Cys Leu His Glu Arg Asn Phe Val Pro Gly Lys Ser Ile Val Glu
640                 645                 650                 655

aat atc atc aac tgc att gag aag agt tac aag tcc atc ttt gtt ttg  2173
Asn Ile Ile Asn Cys Ile Glu Lys Ser Tyr Lys Ser Ile Phe Val Leu
                660                 665                 670

tct ccc aac ttt gtc cag agt gag tgg tgc cat tac gaa ctc tat ttt  2221
Ser Pro Asn Phe Val Gln Ser Glu Trp Cys His Tyr Glu Leu Tyr Phe
            675                 680                 685

gcc cat cac aat ctc ttt cat gaa gga tct aat aac tta atc ctc atc  2269
Ala His His Asn Leu Phe His Glu Gly Ser Asn Asn Leu Ile Leu Ile
        690                 695                 700

tta ctg gaa ccc att cca cag aac agc att ccc aac aag tac cac aag  2317
Leu Leu Glu Pro Ile Pro Gln Asn Ser Ile Pro Asn Lys Tyr His Lys
    705                 710                 715

ctg aag gct ctc atg acg cag cgg act tat ttg cag tgg ccc aag gag  2365
Leu Lys Ala Leu Met Thr Gln Arg Thr Tyr Leu Gln Trp Pro Lys Glu
720                 725                 730                 735

aaa agc aaa cgt ggg ctc ttt tgg gct aac att aga gcc gct ttt aat  2413
Lys Ser Lys Arg Gly Leu Phe Trp Ala Asn Ile Arg Ala Ala Phe Asn
                740                 745                 750
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DX0724XK

atg aaa tta aca cta gtc act gaa aac aat gat gtg aaa tct 2455
Met Lys Leu Thr Leu Val Thr Glu Asn Asn Asp Val Lys Ser
755 760 765

taaaaaaatt taggaaattc aacttaagaa accattattt acttggatga tggtgaatag 2515

tacagtcgta agtnactgtc tggaggtgcc tccattatcc tcatgccttc aggaaagact 2575

taacaaaaac aatgtttcat ctggggaact gagctaggcg gtgaggttag cctgccagtt 2635

agagacagcc cagtctcttc tggtttaatc attatgtttc aaattgaaac agtctctttt 2695

gagtaaatgc tcagtttttc agctcctctc cactctgctt tcccaaatgg attctgttgg 2755

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Glu Phe Ala Val Asp Lys Ser Lys Arg Gly Leu Ile His Val Pro Lys
5 10 15

Asp Leu Pro Leu Lys Thr Lys Val Leu Asp Met Ser Gln Asn Tyr Ile
20 25 30

Ala Glu Leu Gln Val Ser Asp Met Ser Phe Leu Ser Glu Leu Thr Val
35 40 45

Leu Arg Leu Ser His Asn Arg Ile Gln Leu Leu Asp Leu Ser Val Phe
50 55 60 65

Lys Phe Asn Gln Asp Leu Glu Tyr Leu Asp Leu Ser His Asn Gln Leu
70 75 80

Gln Lys Ile Ser Cys His Pro Ile Val Ser Phe Arg His Leu Asp Leu
85 90 95

Ser Phe Asn Asp Phe Lys Ala Leu Pro Ile Cys Lys Glu Phe Gly Asn
100 105 110

Leu Ser Gln Leu Asn Phe Leu Gly Leu Ser Ala Met Lys Leu Gln Lys
115 120 125

Leu Asp Leu Leu Pro Ile Ala His Leu His Leu Ser Tyr Ile Leu Leu

150 155 160

Ile Leu Asn Ala Lys Thr Leu His Leu Val Phe His Pro Thr Ser Leu
165 170 175

Phe Ala Ile Gln Val Asn Ile Ser Val Asn Thr Leu Gly Cys Leu Gln
180 185 190

Leu Thr Asn Ile Lys Leu Asn Asp Asp Asn Cys Gln Val Phe Ile Lys
195 200 205

Phe Leu Ser Glu Leu Thr Arg Gly Pro Thr Leu Leu Asn Phe Thr Leu
210 215 220 225

Asn His Ile Glu Thr Thr Trp Lys Cys Leu Val Arg Val Phe Gln Phe
230 235 240

Leu Trp Pro Lys Pro Val Glu Tyr Leu Asn Ile Tyr Asn Leu Thr Ile
245 250 255

Ile Glu Ser Ile Arg Glu Glu Asp Phe Thr Tyr Ser Lys Thr Thr Leu
260 265 270

Lys Ala Leu Thr Ile Glu His Ile Thr Asn Gln Val Phe Leu Phe Ser
275 280 285

Gln Thr Ala Leu Tyr Thr Val Phe Ser Glu Met Asn Ile Met Met Leu
290 295 300 305

Thr Ile Ser Asp Thr Pro Phe Ile His Met Leu Cys Pro His Ala Pro
310 315 320

Ser Thr Phe Lys Phe Leu Asn Phe Thr Gln Asn Val Phe Thr Asp Ser
325 330 335

Ile Phe Glu Lys Cys Ser Thr Leu Val Lys Leu Glu Thr Leu Ile Leu
340 345 350

Gln Lys Asn Gly Leu Lys Asp Leu Phe Lys Val Gly Leu Met Thr Lys
355 360 365

Asp Met Pro Ser Leu Glu Ile Leu Asp Val Ser Trp Asn Ser Leu Glu
370 375 380 385

Ser Gly Arg His Lys Glu Asn Cys Thr Trp Val Glu Ser Ile Val Val
390 395 400

Leu Asn Leu Ser Ser Asn Met Leu Thr Asp Ser Val Phe Arg Cys Leu
405 410 415

Pro Pro Arg Ile Lys Val Leu Asp Leu His Ser Asn Lys Ile Lys Ser
420 425 430

Val Pro Lys Gln Val Val Lys Leu Glu Ala Leu Gln Glu Leu Asn Val
435 440 445

Ala Phe Asn Ser Leu Thr Asp Leu Pro Gly Cys Gly Ser Phe Ser Ser
450 455 460 465

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Leu Ser Val Leu Ile Ile Asp His Asn Ser Val Ser His Pro Ser Ala
                470                 475                 480

Asp Phe Phe Gln Ser Cys Gln Lys Met Arg Ser Ile Lys Ala Gly Asp
            485                 490                 495

Asn Pro Phe Gln Cys Thr Cys Glu Leu Arg Glu Phe Val Lys Asn Ile
        500                 505                 510

Asp Gln Val Ser Ser Glu Val Leu Glu Gly Trp Pro Asp Ser Tyr Lys
    515                 520                 525

Cys Asp Tyr Pro Glu Ser Tyr Arg Gly Ser Pro Leu Lys Asp Phe His
530                 535                 540                 545

Met Ser Glu Leu Ser Cys Asn Ile Thr Leu Leu Ile Val Thr Ile Gly
                550                 555                 560

Ala Thr Met Leu Val Leu Ala Val Thr Val Thr Ser Leu Cys Ile Tyr
            565                 570                 575

Leu Asp Leu Pro Trp Tyr Leu Arg Met Val Cys Gln Trp Thr Gln Thr
        580                 585                 590

Arg Arg Arg Ala Arg Asn Ile Pro Leu Glu Glu Leu Gln Arg Asn Leu
    595                 600                 605

Gln Phe His Ala Phe Ile Ser Tyr Ser Glu His Asp Ser Ala Trp Val
610                 615                 620                 625

Lys Ser Glu Leu Val Pro Tyr Leu Glu Lys Glu Asp Ile Gln Ile Cys
                630                 635                 640

Leu His Glu Arg Asn Phe Val Pro Gly Lys Ser Ile Val Glu Asn Ile
            645                 650                 655

Ile Asn Cys Ile Glu Lys Ser Tyr Lys Ser Ile Phe Val Leu Ser Pro
        660                 665                 670

Asn Phe Val Gln Ser Glu Trp Cys His Tyr Glu Leu Tyr Phe Ala His
    675                 680                 685

His Asn Leu Phe His Glu Gly Ser Asn Asn Leu Ile Leu Ile Leu Leu
690                 695                 700                 705

Glu Pro Ile Pro Gln Asn Ser Ile Pro Asn Lys Tyr His Lys Leu Lys
                710                 715                 720

Ala Leu Met Thr Gln Arg Thr Tyr Leu Gln Trp Pro Lys Glu Lys Ser
            725                 730                 735

Lys Arg Gly Leu Phe Trp Ala Asn Ile Arg Ala Ala Phe Asn Met Lys
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cac aca gcc ctc cca ccc cca cag ggt ttc tgc cgc agc gcc ctg cac   96
His Thr Ala Leu Pro Pro Pro Gln Gly Phe Cys Arg Ser Ala Leu His
        -30                 -25                 -20

ccg ctg tct ctc ctg gtg cag gcc atc atg ctg gcc atg acc ctg gcc   144
Pro Leu Ser Leu Leu Val Gln Ala Ile Met Leu Ala Met Thr Leu Ala
    -15                 -10                 -5                  -1

ctg ggt acc ttg cct gcc ttc cta ccc tgt gag ctc cag ccc cac ggc   192
Leu Gly Thr Leu Pro Ala Phe Leu Pro Cys Glu Leu Gln Pro His Gly
  1               5                   10                  15

ctg gtg aac tgc aac tgg ctg ttc ctg aag tct gtg ccc cac ttc tcc   240
Leu Val Asn Cys Asn Trp Leu Phe Leu Lys Ser Val Pro His Phe Ser
            20                  25                  30

atg gca gca ccc cgt ggc aat gtc acc agc ctt tcc ttg tcc tcc aac   288
Met Ala Ala Pro Arg Gly Asn Val Thr Ser Leu Ser Leu Ser Ser Asn
        35                  40                  45

cgc atc cac cac ctc cat gat tct gac ttt gcc cac ctg ccc agc ctg   336
Arg Ile His His Leu His Asp Ser Asp Phe Ala His Leu Pro Ser Leu
    50                  55                  60

cgg cat ctc aac ctc aag tgg aac tgc ccg ccg gtt ggc ctc agc ccc   384
Arg His Leu Asn Leu Lys Trp Asn Cys Pro Pro Val Gly Leu Ser Pro
65                  70                  75                  80

atg cac ttc ccc tgc cac atg acc atc gag ccc agc acc ttc ttg gct   432
Met His Phe Pro Cys His Met Thr Ile Glu Pro Ser Thr Phe Leu Ala
                85                  90                  95

gtg ccc acc ctg gaa gag cta aac ctg agc tac aac aac atc atg act   480
Val Pro Thr Leu Glu Glu Leu Asn Leu Ser Tyr Asn Asn Ile Met Thr
            100                 105                 110
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gtg cct gcg ctg ccc aaa tcc ctc ata tcc ctg tcc ctc agc cat acc   528
Val Pro Ala Leu Pro Lys Ser Leu Ile Ser Leu Ser Leu Ser His Thr
        115                 120                 125

aac atc ctg atg cta gac tct gcc agc ctc gcc ggc ctg cat gcc ctg   576
Asn Ile Leu Met Leu Asp Ser Ala Ser Leu Ala Gly Leu His Ala Leu
    130                 135                 140

cgc ttc cta ttc atg gac ggc aac tgt tat tac aag aac ccc tgc agg   624
Arg Phe Leu Phe Met Asp Gly Asn Cys Tyr Tyr Lys Asn Pro Cys Arg
145                 150                 155                 160

cag gca ctg gag gtg gcc ccg ggt gcc ctc ctt ggc ctg ggc aac ctc   672
Gln Ala Leu Glu Val Ala Pro Gly Ala Leu Leu Gly Leu Gly Asn Leu
                165                 170                 175

acc cac ctg tca ctc aag tac aac aac ctc act gtg gtg ccc cgc aac   720
Thr His Leu Ser Leu Lys Tyr Asn Asn Leu Thr Val Val Pro Arg Asn
            180                 185                 190

ctg cct tcc agc ctg gag tat ctg ctg ttg tcc tac aac cgc atc gtc   768
Leu Pro Ser Ser Leu Glu Tyr Leu Leu Leu Ser Tyr Asn Arg Ile Val
        195                 200                 205

aaa ctg gcg cct gag gac ctg gcc aat ctg acc gcc ctg cgt gtg ctc   816
Lys Leu Ala Pro Glu Asp Leu Ala Asn Leu Thr Ala Leu Arg Val Leu
    210                 215                 220

gat gtg ggc gga aat tgc cgc cgc tgc gac cac gct ccc aac ccc tgc   864
Asp Val Gly Gly Asn Cys Arg Arg Cys Asp His Ala Pro Asn Pro Cys
225                 230                 235                 240

atg gag tgc cct cgt cac ttc ccc cag cta cat ccc gat acc ttc agc   912
Met Glu Cys Pro Arg His Phe Pro Gln Leu His Pro Asp Thr Phe Ser
                245                 250                 255

cac ctg agc cgt ctt gaa ggc ctg gtg ttg aag gac agt tct ctc tcc   960
His Leu Ser Arg Leu Glu Gly Leu Val Leu Lys Asp Ser Ser Leu Ser
            260                 265                 270

tgg ctg aat gcc agt tgg ttc cgt ggg ctg gga aac ctc cga gtg ctg   1008
Trp Leu Asn Ala Ser Trp Phe Arg Gly Leu Gly Asn Leu Arg Val Leu
        275                 280                 285

gac ctg agt gag aac ttc ctc tac aaa tgc atc act aaa acc aag gcc   1056
Asp Leu Ser Glu Asn Phe Leu Tyr Lys Cys Ile Thr Lys Thr Lys Ala
    290                 295                 300

ttc cag ggc cta aca cag ctg cgc aag ctt aac ctg tcc ttc aat tac   1104
Phe Gln Gly Leu Thr Gln Leu Arg Lys Leu Asn Leu Ser Phe Asn Tyr
305                 310                 315                 320

caa aag agg gtg tcc ttt gcc cac ctg tct ctg gcc cct tcc ttc ggg   1152
Gln Lys Arg Val Ser Phe Ala His Leu Ser Leu Ala Pro Ser Phe Gly
                325                 330                 335

agc ctg gtc gcc ctg aag gag ctg gac atg cac ggc atc ttc ttc cgc   1200
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Ser Leu Val Ala Leu Lys Glu Leu Asp Met His Gly Ile Phe Phe Arg
            340                 345                 350

tca ctc gat gag acc acg ctc cgg cca ctg gcc cgc ctg ccc atg ctc  1248
Ser Leu Asp Glu Thr Thr Leu Arg Pro Leu Ala Arg Leu Pro Met Leu
        355                 360                 365

cag act ctg cgt ctg cag atg aac ttc atc aac cag gcc cag ctc ggc  1296
Gln Thr Leu Arg Leu Gln Met Asn Phe Ile Asn Gln Ala Gln Leu Gly
    370                 375                 380

atc ttc agg gcc ttc cct ggc ctg cgc tac gtg gac ctg tcg gac aac  1344
Ile Phe Arg Ala Phe Pro Gly Leu Arg Tyr Val Asp Leu Ser Asp Asn
385                 390                 395                 400

cgc atc agc gga gct tcg gag ctg aca gcc acc atg ggg gag gca gat  1392
Arg Ile Ser Gly Ala Ser Glu Leu Thr Ala Thr Met Gly Glu Ala Asp
                405                 410                 415

gga ggg gag aag gtc tgg ctg cag cct ggg gac ctt gct ccg gcc cca  1440
Gly Gly Glu Lys Val Trp Leu Gln Pro Gly Asp Leu Ala Pro Ala Pro
            420                 425                 430

gtg gac act ccc agc tct gaa gac ttc agg ccc aac tgc agc acc ctc  1488
Val Asp Thr Pro Ser Ser Glu Asp Phe Arg Pro Asn Cys Ser Thr Leu
        435                 440                 445

aac ttc acc ttg gat ctg tca cgg aac aac ctg gtg acc gtg cag ccg  1536
Asn Phe Thr Leu Asp Leu Ser Arg Asn Asn Leu Val Thr Val Gln Pro
    450                 455                 460

gag atg ttt gcc cag ctc tcg cac ctg cag tgc ctg cgc ctg agc cac  1584
Glu Met Phe Ala Gln Leu Ser His Leu Gln Cys Leu Arg Leu Ser His
465                 470                 475                 480

aac tgc atc tcg cag gca gtc aat ggc tcc cag ttc ctg ccg ctg acc  1632
Asn Cys Ile Ser Gln Ala Val Asn Gly Ser Gln Phe Leu Pro Leu Thr
                485                 490                 495

ggt ctg cag gtg cta gac ctg tcc cac aat aag ctg gac ctc tac cac  1680
Gly Leu Gln Val Leu Asp Leu Ser His Asn Lys Leu Asp Leu Tyr His
            500                 505                 510

gag cac tca ttc acg gag cta cca cga ctg gag gcc ctg gac ctc agc  1728
Glu His Ser Phe Thr Glu Leu Pro Arg Leu Glu Ala Leu Asp Leu Ser
        515                 520                 525

tac aac agc cag ccc ttt ggc atg cag ggc gtg ggc cac aac ttc agc  1776
Tyr Asn Ser Gln Pro Phe Gly Met Gln Gly Val Gly His Asn Phe Ser
    530                 535                 540

ttc gtg gct cac ctg cgc acc ctg cgc cac ctc agc ctg gcc cac aac  1824
Phe Val Ala His Leu Arg Thr Leu Arg His Leu Ser Leu Ala His Asn
545                 550                 555                 560

aac atc cac agc caa gtg tcc cag cag ctc tgc agt acg tcg ctg cgg  1872
Asn Ile His Ser Gln Val Ser Gln Gln Leu Cys Ser Thr Ser Leu Arg
                565                 570                 575
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gcc ctg gac ttc agc ggc aat gca ctg ggc cat atg tgg gcc gag gga 1920
Ala Leu Asp Phe Ser Gly Asn Ala Leu Gly His Met Trp Ala Glu Gly
            580                 585                 590

gac ctc tat ctg cac ttc ttc caa ggc ctg agc ggt ttg atc tgg ctg 1968
Asp Leu Tyr Leu His Phe Phe Gln Gly Leu Ser Gly Leu Ile Trp Leu
        595                 600                 605

gac ttg tcc cag aac cgc ctg cac acc ctc ctg ccc caa acc ctg cgc 2016
Asp Leu Ser Gln Asn Arg Leu His Thr Leu Leu Pro Gln Thr Leu Arg
    610                 615                 620

aac ctc ccc aag agc cta cag gtg ctg cgt ctc cgt gac aat tac ctg 2064
Asn Leu Pro Lys Ser Leu Gln Val Leu Arg Leu Arg Asp Asn Tyr Leu
625                 630                 635                 640

gcc ttc ttt aag tgg tgg agc ctc cac ttc ctg ccc aaa ctg gaa gtc 2112
Ala Phe Phe Lys Trp Trp Ser Leu His Phe Leu Pro Lys Leu Glu Val
                645                 650                 655

ctc gac ctg gca gga aac cag ctg aag gcc ctg acc aat ggc agc ctg 2160
Leu Asp Leu Ala Gly Asn Gln Leu Lys Ala Leu Thr Asn Gly Ser Leu
            660                 665                 670

cct gct ggc acc cgg ctc cgg agg ctg gat gtc agc tgc aac agc atc 2208
Pro Ala Gly Thr Arg Leu Arg Arg Leu Asp Val Ser Cys Asn Ser Ile
        675                 680                 685

agc ttc gtg gcc ccc ggc ttc ttt tcc aag gcc aag gag ctg cga gag 2256
Ser Phe Val Ala Pro Gly Phe Phe Ser Lys Ala Lys Glu Leu Arg Glu
    690                 695                 700

ctc aac ctt agc gcc aac gcc ctc aag aca gtg gac cac tcc tgg ttt 2304
Leu Asn Leu Ser Ala Asn Ala Leu Lys Thr Val Asp His Ser Trp Phe
705                 710                 715                 720

ggg ccc ctg gcg agt gcc ctg caa ata cta gat gta agc gcc aac cct 2352
Gly Pro Leu Ala Ser Ala Leu Gln Ile Leu Asp Val Ser Ala Asn Pro
                725                 730                 735

ctg cac tgc gcc tgt ggg gcg gcc ttt atg gac ttc ctg ctg gag gtg 2400
Leu His Cys Ala Cys Gly Ala Ala Phe Met Asp Phe Leu Leu Glu Val
            740                 745                 750

cag gct gcc gtg ccc ggt ctg ccc agc cgg gtg aag tgt ggc agt ccg 2448
Gln Ala Ala Val Pro Gly Leu Pro Ser Arg Val Lys Cys Gly Ser Pro
        755                 760                 765

ggc cag ctc cag ggc ctc agc atc ttt gca cag gac ctg cgc ctc tgc 2496
Gly Gln Leu Gln Gly Leu Ser Ile Phe Ala Gln Asp Leu Arg Leu Cys
    770                 775                 780

ctg gat gag gcc ctc tcc tgg gac tgt ttc gcc ctc tcg ctg ctg gct 2544
Leu Asp Glu Ala Leu Ser Trp Asp Cys Phe Ala Leu Ser Leu Leu Ala
785                 790                 795                 800

gtg gct ctg ggc ctg ggt gtg ccc atg ctg cat cac ctc tgt ggc tgg 2592
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Val Ala Leu Gly Leu Gly Val Pro Met Leu His His Leu Cys Gly Trp
                805                 810                 815

gac ctc tgg tac tgc ttc cac ctg tgc ctg gcc tgg ctt ccc tgg cgg   2640
Asp Leu Trp Tyr Cys Phe His Leu Cys Leu Ala Trp Leu Pro Trp Arg
            820                 825                 830

ggg cgg caa agt ggg cga gat gag gat gcc ctg ccc tac gat gcc ttc   2688
Gly Arg Gln Ser Gly Arg Asp Glu Asp Ala Leu Pro Tyr Asp Ala Phe
        835                 840                 845

gtg gtc ttc gac aaa acg cag agc gca gtg gca gac tgg gtg tac aac   2736
Val Val Phe Asp Lys Thr Gln Ser Ala Val Ala Asp Trp Val Tyr Asn
    850                 855                 860

gag ctt cgg ggg cag ctg gag gag tgc cgt ggg cgc tgg gca ctc cgc   2784
Glu Leu Arg Gly Gln Leu Glu Glu Cys Arg Gly Arg Trp Ala Leu Arg
865                 870                 875                 880

ctg tgc ctg gag gaa cgc gac tgg ctg cct ggc aaa acc ctc ttt gag   2832
Leu Cys Leu Glu Glu Arg Asp Trp Leu Pro Gly Lys Thr Leu Phe Glu
                885                 890                 895

aac ctg tgg gcc tcg gtc tat ggc agc cgc aag acg ctg ttt gtg ctg   2880
Asn Leu Trp Ala Ser Val Tyr Gly Ser Arg Lys Thr Leu Phe Val Leu
            900                 905                 910

gcc cac acg gac cgg gtc agt ggt ctc ttg cgc gcc agc ttc ctg ctg   2928
Ala His Thr Asp Arg Val Ser Gly Leu Leu Arg Ala Ser Phe Leu Leu
        915                 920                 925

gcc cag cag cgc ctg ctg gag gac cgc aag gac gtc gtg gtg ctg gtg   2976
Ala Gln Gln Arg Leu Leu Glu Asp Arg Lys Asp Val Val Val Leu Val
    930                 935                 940

atc ctg agc cct gac ggc cgc cgc tcc cgc tat gtg cgg ctg cgc cag   3024
Ile Leu Ser Pro Asp Gly Arg Arg Ser Arg Tyr Val Arg Leu Arg Gln
945                 950                 955                 960

cgc ctc tgc cgc cag agt gtc ctc ctc tgg ccc cac cag ccc agt ggt   3072
Arg Leu Cys Arg Gln Ser Val Leu Leu Trp Pro His Gln Pro Ser Gly
                965                 970                 975

cag cgc agc ttc tgg gcc cag ctg ggc atg gcc ctg acc agg gac aac   3120
Gln Arg Ser Phe Trp Ala Gln Leu Gly Met Ala Leu Thr Arg Asp Asn
            980                 985                 990

cac cac ttc tat aac cgg aac ttc tgc cag gga ccc acg gcc gaa tag   3168
His His Phe Tyr Asn Arg Asn Phe Cys Gln Gly Pro Thr Ala Glu
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His Thr Ala Leu Pro Pro Pro Gln Gly Phe Cys Arg Ser Ala Leu His
          -30                 -25                 -20

Pro Leu Ser Leu Leu Val Gln Ala Ile Met Leu Ala Met Thr Leu Ala
     -15                 -10                  -5                  -1

Leu Gly Thr Leu Pro Ala Phe Leu Pro Cys Glu Leu Gln Pro His Gly
  1                   5                  10                  15

Leu Val Asn Cys Asn Trp Leu Phe Leu Lys Ser Val Pro His Phe Ser
             20                  25                  30

Met Ala Ala Pro Arg Gly Asn Val Thr Ser Leu Ser Leu Ser Ser Asn
          35                  40                  45

Arg Ile His His Leu His Asp Ser Asp Phe Ala His Leu Pro Ser Leu
      50                  55                  60

Arg His Leu Asn Leu Lys Trp Asn Cys Pro Pro Val Gly Leu Ser Pro
  65                  70                  75                  80

Met His Phe Pro Cys His Met Thr Ile Glu Pro Ser Thr Phe Leu Ala
                 85                  90                  95

Val Pro Thr Leu Glu Glu Leu Asn Leu Ser Tyr Asn Asn Ile Met Thr
            100                 105                 110

Val Pro Ala Leu Pro Lys Ser Leu Ile Ser Leu Ser Leu Ser His Thr
        115                 120                 125

Asn Ile Leu Met Leu Asp Ser Ala Ser Leu Ala Gly Leu His Ala Leu
    130                 135                 140

Arg Phe Leu Phe Met Asp Gly Asn Cys Tyr Tyr Lys Asn Pro Cys Arg
145                 150                 155                 160

Gln Ala Leu Glu Val Ala Pro Gly Ala Leu Leu Gly Leu Gly Asn Leu
                165                 170                 175

Thr His Leu Ser Leu Lys Tyr Asn Asn Leu Thr Val Val Pro Arg Asn
            180                 185                 190

Leu Pro Ser Ser Leu Glu Tyr Leu Leu Leu Ser Tyr Asn Arg Ile Val
        195                 200                 205

Lys Leu Ala Pro Glu Asp Leu Ala Asn Leu Thr Ala Leu Arg Val Leu
    210                 215                 220

Asp Val Gly Gly Asn Cys Arg Arg Cys Asp His Ala Pro Asn Pro Cys
225                 230                 235                 240

Met Glu Cys Pro Arg His Phe Pro Gln Leu His Pro Asp Thr Phe Ser
                245                 250                 255

His Leu Ser Arg Leu Glu Gly Leu Val Leu Lys Asp Ser Ser Leu Ser
```

260 265 270

Trp Leu Asn Ala Ser Trp Phe Arg Gly Leu Gly Asn Leu Arg Val Leu
275 280 285

Asp Leu Ser Glu Asn Phe Leu Tyr Lys Cys Ile Thr Lys Thr Lys Ala
290 295 300

Phe Gln Gly Leu Thr Gln Leu Arg Lys Leu Asn Leu Ser Phe Asn Tyr
305 310 315 320

Gln Lys Arg Val Ser Phe Ala His Leu Ser Leu Ala Pro Ser Phe Gly
325 330 335

Ser Leu Val Ala Leu Lys Glu Leu Asp Met His Gly Ile Phe Phe Arg
340 345 350

Ser Leu Asp Glu Thr Thr Leu Arg Pro Leu Ala Arg Leu Pro Met Leu
355 360 365

Gln Thr Leu Arg Leu Gln Met Asn Phe Ile Asn Gln Ala Gln Leu Gly
370 375 380

Ile Phe Arg Ala Phe Pro Gly Leu Arg Tyr Val Asp Leu Ser Asp Asn
385 390 395 400

Arg Ile Ser Gly Ala Ser Glu Leu Thr Ala Thr Met Gly Glu Ala Asp
405 410 415

Gly Gly Glu Lys Val Trp Leu Gln Pro Gly Asp Leu Ala Pro Ala Pro
420 425 430

Val Asp Thr Pro Ser Ser Glu Asp Phe Arg Pro Asn Cys Ser Thr Leu
435 440 445

Asn Phe Thr Leu Asp Leu Ser Arg Asn Asn Leu Val Thr Val Gln Pro
450 455 460

Glu Met Phe Ala Gln Leu Ser His Leu Gln Cys Leu Arg Leu Ser His
465 470 475 480

Asn Cys Ile Ser Gln Ala Val Asn Gly Ser Gln Phe Leu Pro Leu Thr
485 490 495

Gly Leu Gln Val Leu Asp Leu Ser His Asn Lys Leu Asp Leu Tyr His
500 505 510

Glu His Ser Phe Thr Glu Leu Pro Arg Leu Glu Ala Leu Asp Leu Ser
515 520 525

Tyr Asn Ser Gln Pro Phe Gly Met Gln Gly Val Gly His Asn Phe Ser
530 535 540

Phe Val Ala His Leu Arg Thr Leu Arg His Leu Ser Leu Ala His Asn
545 550 555 560

Asn Ile His Ser Gln Val Ser Gln Gln Leu Cys Ser Thr Ser Leu Arg
565 570 575

```
Ala Leu Asp Phe Ser Gly Asn Ala Leu Gly His Met Trp Ala Glu Gly
            580                 585                 590

Asp Leu Tyr Leu His Phe Phe Gln Gly Leu Ser Gly Leu Ile Trp Leu
        595                 600                 605

Asp Leu Ser Gln Asn Arg Leu His Thr Leu Leu Pro Gln Thr Leu Arg
    610                 615                 620

Asn Leu Pro Lys Ser Leu Gln Val Leu Arg Leu Arg Asp Asn Tyr Leu
625                 630                 635                 640

Ala Phe Phe Lys Trp Trp Ser Leu His Phe Leu Pro Lys Leu Glu Val
                645                 650                 655

Leu Asp Leu Ala Gly Asn Gln Leu Lys Ala Leu Thr Asn Gly Ser Leu
            660                 665                 670

Pro Ala Gly Thr Arg Leu Arg Arg Leu Asp Val Ser Cys Asn Ser Ile
        675                 680                 685

Ser Phe Val Ala Pro Gly Phe Phe Ser Lys Ala Lys Glu Leu Arg Glu
    690                 695                 700

Leu Asn Leu Ser Ala Asn Ala Leu Lys Thr Val Asp His Ser Trp Phe
705                 710                 715                 720

Gly Pro Leu Ala Ser Ala Leu Gln Ile Leu Asp Val Ser Ala Asn Pro
                725                 730                 735

Leu His Cys Ala Cys Gly Ala Ala Phe Met Asp Phe Leu Leu Glu Val
            740                 745                 750

Gln Ala Ala Val Pro Gly Leu Pro Ser Arg Val Lys Cys Gly Ser Pro
        755                 760                 765

Gly Gln Leu Gln Gly Leu Ser Ile Phe Ala Gln Asp Leu Arg Leu Cys
    770                 775                 780

Leu Asp Glu Ala Leu Ser Trp Asp Cys Phe Ala Leu Ser Leu Leu Ala
785                 790                 795                 800

Val Ala Leu Gly Leu Gly Val Pro Met Leu His His Leu Cys Gly Trp
                805                 810                 815

Asp Leu Trp Tyr Cys Phe His Leu Cys Leu Ala Trp Leu Pro Trp Arg
            820                 825                 830

Gly Arg Gln Ser Gly Arg Asp Glu Asp Ala Leu Pro Tyr Asp Ala Phe
        835                 840                 845

Val Val Phe Asp Lys Thr Gln Ser Ala Val Ala Asp Trp Val Tyr Asn
    850                 855                 860

Glu Leu Arg Gly Gln Leu Glu Glu Cys Arg Gly Arg Trp Ala Leu Arg
865                 870                 875                 880
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Leu Cys Leu Glu Glu Arg Asp Trp Leu Pro Gly Lys Thr Leu Phe Glu
885 890 895

Asn Leu Trp Ala Ser Val Tyr Gly Ser Arg Lys Thr Leu Phe Val Leu
900 905 910

Ala His Thr Asp Arg Val Ser Gly Leu Leu Arg Ala Ser Phe Leu Leu
915 920 925

Ala Gln Gln Arg Leu Leu Glu Asp Arg Lys Asp Val Val Val Leu Val
930 935 940

Ile Leu Ser Pro Asp Gly Arg Arg Ser Arg Tyr Val Arg Leu Arg Gln
945 950 955 960

Arg Leu Cys Arg Gln Ser Val Leu Leu Trp Pro His Gln Pro Ser Gly
965 970 975

Gln Arg Ser Phe Trp Ala Gln Leu Gly Met Ala Leu Thr Arg Asp Asn
980 985 990

His His Phe Tyr Asn Arg Asn Phe Cys Gln Gly Pro Thr Ala Glu
995 1000 1005

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Mus musculus

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ctg gca agt tcc ttt aag aac ctg gtg tca ctg cag gag ctg aac atg 96
Leu Ala Ser Ser Phe Lys Asn Leu Val Ser Leu Gln Glu Leu Asn Met
20 25 30

aac ggc atc ttc ttc cgc ttg ctc aac aag tac acg ctc aga tgg ctg 144
Asn Gly Ile Phe Phe Arg Leu Leu Asn Lys Tyr Thr Leu Arg Trp Leu
35 40 45

gcc gat ctg ccc aaa ctc cac act ctg cat ctt caa atg aac ttc atc 192
Ala Asp Leu Pro Lys Leu His Thr Leu His Leu Gln Met Asn Phe Ile
50 55 60

aac cag gca cag ctc agc atc ttt ggt acc ttc cga gcc ctt cgc ttt 240
Asn Gln Ala Gln Leu Ser Ile Phe Gly Thr Phe Arg Ala Leu Arg Phe
65 70 75 80

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gtg gac ttg tca gac aat cgc atc agt ggg cct tca acg ctg tca gaa   288
Val Asp Leu Ser Asp Asn Arg Ile Ser Gly Pro Ser Thr Leu Ser Glu
                 85                  90                  95

gcc acc cct gaa gag gca gat gat gca gag cag gag gag ctg ttg tct   336
Ala Thr Pro Glu Glu Ala Asp Asp Ala Glu Gln Glu Glu Leu Leu Ser
            100                 105                 110

gcg gat cct cac cca gct ccg ctg agc acc cct gct tct aag aac ttc   384
Ala Asp Pro His Pro Ala Pro Leu Ser Thr Pro Ala Ser Lys Asn Phe
        115                 120                 125

atg gac agg tgt aag aac ttc aag ttc aac atg gac ctg tct cgg aac   432
Met Asp Arg Cys Lys Asn Phe Lys Phe Asn Met Asp Leu Ser Arg Asn
    130                 135                 140

aac ctg gtg act atc aca gca gag atg ttt gta aat ctc tca cgc ctc   480
Asn Leu Val Thr Ile Thr Ala Glu Met Phe Val Asn Leu Ser Arg Leu
145                 150                 155                 160

cag tgt ctt agc ctg agc cac aac tca att gca cag gct gtc aat ggc   528
Gln Cys Leu Ser Leu Ser His Asn Ser Ile Ala Gln Ala Val Asn Gly
                165                 170                 175

tct cag ttc ctg ccg ctg acc ggt ctg cag gtg cta gac ctg tcc cac   576
Ser Gln Phe Leu Pro Leu Thr Gly Leu Gln Val Leu Asp Leu Ser His
            180                 185                 190

aat aag ctg gac ctc tac cac gag cac tca ttc acg gag cta cca cga   624
Asn Lys Leu Asp Leu Tyr His Glu His Ser Phe Thr Glu Leu Pro Arg
        195                 200                 205

ctg gag gcc ctg gac ctc agc tac aac agc cag ccc ttt agc atg aag   672
Leu Glu Ala Leu Asp Leu Ser Tyr Asn Ser Gln Pro Phe Ser Met Lys
    210                 215                 220

ggt ata ggc cac aat ttc agt ttt gtg acc cat ctg tcc atg cta cag   720
Gly Ile Gly His Asn Phe Ser Phe Val Thr His Leu Ser Met Leu Gln
225                 230                 235                 240

agc ctt agc ctg gca cac aat gac att cat acc cgt gtg tcc tca cat   768
Ser Leu Ser Leu Ala His Asn Asp Ile His Thr Arg Val Ser Ser His
                245                 250                 255

ctc aac agc aac tca gtg agg ttt ctt gac ttc agc ggc aac ggt atg   816
Leu Asn Ser Asn Ser Val Arg Phe Leu Asp Phe Ser Gly Asn Gly Met
            260                 265                 270

ggc cgc atg tgg gat gag ggg ggc ctt tat ctc cat ttc ttc caa ggc   864
Gly Arg Met Trp Asp Glu Gly Gly Leu Tyr Leu His Phe Phe Gln Gly
        275                 280                 285

ctg agt ggc gtg ctg aag ctg gac ctg tct caa aat aac ctg cat atc   912
Leu Ser Gly Val Leu Lys Leu Asp Leu Ser Gln Asn Asn Leu His Ile
    290                 295                 300

ctc cgg ccc cag aac ctt gac aac ctc ccc aag agc ctg aag ctg ctg   960
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Leu Arg Pro Gln Asn Leu Asp Asn Leu Pro Lys Ser Leu Lys Leu Leu
305                 310                 315                 320

agc ctc cga gac aac tac cta tct ttc ttt aac tgg acc agt ctg tcc 1008
Ser Leu Arg Asp Asn Tyr Leu Ser Phe Phe Asn Trp Thr Ser Leu Ser
                325                 330                 335

ttc cta ccc aac ctg gaa gtc cta gac ctg gca ggc aac cag cta aag 1056
Phe Leu Pro Asn Leu Glu Val Leu Asp Leu Ala Gly Asn Gln Leu Lys
            340                 345                 350

gcc ctg acc aat ggc acc ctg cct aat ggc acc ctc ctc cag aaa ctc 1104
Ala Leu Thr Asn Gly Thr Leu Pro Asn Gly Thr Leu Leu Gln Lys Leu
        355                 360                 365

gat gtc agt agc aac agt atc gtc tct gtg gcc ccc ggc ttc ttt tcc 1152
Asp Val Ser Ser Asn Ser Ile Val Ser Val Ala Pro Gly Phe Phe Ser
    370                 375                 380

aag gcc aag gag ctg cga gag ctc aac ctt agc gcc aac gcc ctc aag 1200
Lys Ala Lys Glu Leu Arg Glu Leu Asn Leu Ser Ala Asn Ala Leu Lys
385                 390                 395                 400

aca gtg gac cac tcc tgg ttt ggg ccc att gtg atg aac ctg aca gtt 1248
Thr Val Asp His Ser Trp Phe Gly Pro Ile Val Met Asn Leu Thr Val
                405                 410                 415

cta gac gtg aga agc aac cct ctg cac tgt gcc tgt ggg gca gcc ttc 1296
Leu Asp Val Arg Ser Asn Pro Leu His Cys Ala Cys Gly Ala Ala Phe
            420                 425                 430

gta gac tta ctg ttg gag gtg cag acc aag gtg cct ggc ctg gct aat 1344
Val Asp Leu Leu Leu Glu Val Gln Thr Lys Val Pro Gly Leu Ala Asn
        435                 440                 445

ggt gtg aag tgt ggc agc ccc ggc cag ctg cag ggc cgt agc atc ttc 1392
Gly Val Lys Cys Gly Ser Pro Gly Gln Leu Gln Gly Arg Ser Ile Phe
    450                 455                 460

gcg cag gac ctg cgg ctg tgc ctg gat gag gtc ctc tct tgg gac tgc 1440
Ala Gln Asp Leu Arg Leu Cys Leu Asp Glu Val Leu Ser Trp Asp Cys
465                 470                 475                 480

ttt ggc ctt tca ctc ttg gct gtg gcc gtg ggc atg gtg gtg cct ata 1488
Phe Gly Leu Ser Leu Leu Ala Val Ala Val Gly Met Val Val Pro Ile
                485                 490                 495

ctg cac cat ctc tgc ggc tgg gac gtc tgg tac tgt ttt cat ctg tgc 1536
Leu His His Leu Cys Gly Trp Asp Val Trp Tyr Cys Phe His Leu Cys
            500                 505                 510

ctg gca tgg cta cct ttg cta gcc cgc agc cga cgc agc gcc caa act 1584
Leu Ala Trp Leu Pro Leu Leu Ala Arg Ser Arg Arg Ser Ala Gln Thr
        515                 520                 525

ctc cct tat gat gcc ttc gtg gtg ttc gat aag gca cag agc gca gtt 1632
Leu Pro Tyr Asp Ala Phe Val Val Phe Asp Lys Ala Gln Ser Ala Val
    530                 535                 540
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gcc gac tgg gtg tat aac gag ctg cgg gtg cgg ctg gag gag cgg cgc   1680
Ala Asp Trp Val Tyr Asn Glu Leu Arg Val Arg Leu Glu Glu Arg Arg
545                 550                 555                 560

ggc cgc tgg gca ctc cgc ctg tgc ctg gag gac cga gat tgg ctg cct   1728
Gly Arg Trp Ala Leu Arg Leu Cys Leu Glu Asp Arg Asp Trp Leu Pro
                565                 570                 575

ggc cag acg ctc ttc gag aac ctc tgg gct tcc atc tat ggg agc cgc   1776
Gly Gln Thr Leu Phe Glu Asn Leu Trp Ala Ser Ile Tyr Gly Ser Arg
            580                 585                 590

aag act cta ttt gtg ctg gcc cac acg gac cgc gtc agt ggc ctc ctg   1824
Lys Thr Leu Phe Val Leu Ala His Thr Asp Arg Val Ser Gly Leu Leu
        595                 600                 605

cgc acc agc ttc ctg ctg gct cag cag cgc ctg ttg gaa gac cgc aag   1872
Arg Thr Ser Phe Leu Leu Ala Gln Gln Arg Leu Leu Glu Asp Arg Lys
    610                 615                 620

gac gtg gtg gtg ttg gtg atc ctg cgt ccg gat gcc cac cgc tcc cgc   1920
Asp Val Val Val Leu Val Ile Leu Arg Pro Asp Ala His Arg Ser Arg
625                 630                 635                 640

tat gtg cga ctg cgc cag cgt ctc tgc cgc cag agt gtg ctc ttc tgg   1968
Tyr Val Arg Leu Arg Gln Arg Leu Cys Arg Gln Ser Val Leu Phe Trp
                645                 650                 655

ccc cag cag ccc aac ggg cag ggg ggc ttc tgg gcc cag ctg agt aca   2016
Pro Gln Gln Pro Asn Gly Gln Gly Gly Phe Trp Ala Gln Leu Ser Thr
            660                 665                 670

gcc ctg act agg gac aac cgc cac ttc tat aac cag aac ttc tgc cgg   2064
Ala Leu Thr Arg Asp Asn Arg His Phe Tyr Asn Gln Asn Phe Cys Arg
        675                 680                 685

gga ccc aca gca gaa tagctcagag caacagctgg aaacagctgc atcttcatgt   2119
Gly Pro Thr Ala Glu
    690

ctggttcccg agttgctctg cctgccttgc tctgtcttac tacaccgcta tttggcaagt 2179

gcgcaatata tgctaccaag ccaccaggcc cacggagcaa aggttggctg taaagggtag 2239

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20 25 30

Asn Gly Ile Phe Phe Arg Leu Leu Asn Lys Tyr Thr Leu Arg Trp Leu
35 40 45

Ala Asp Leu Pro Lys Leu His Thr Leu His Leu Gln Met Asn Phe Ile
50 55 60

Asn Gln Ala Gln Leu Ser Ile Phe Gly Thr Phe Arg Ala Leu Arg Phe
65 70 75 80

Val Asp Leu Ser Asp Asn Arg Ile Ser Gly Pro Ser Thr Leu Ser Glu
85 90 95

Ala Thr Pro Glu Glu Ala Asp Asp Ala Glu Gln Glu Glu Leu Leu Ser
100 105 110

Ala Asp Pro His Pro Ala Pro Leu Ser Thr Pro Ala Ser Lys Asn Phe
115 120 125

Met Asp Arg Cys Lys Asn Phe Lys Phe Asn Met Asp Leu Ser Arg Asn
130 135 140

Asn Leu Val Thr Ile Thr Ala Glu Met Phe Val Asn Leu Ser Arg Leu
145 150 160

Gln Cys Leu Ser Leu Ser His Asn Ser Ile Ala Gln Ala Val Asn Gly
165 170 175

Ser Gln Phe Leu Pro Leu Thr Gly Leu Gln Val Leu Asp Leu Ser His
180 185 190

Asn Lys Leu Asp Leu Tyr His Glu His Ser Phe Thr Glu Leu Pro Arg
195 200 205

Leu Glu Ala Leu Asp Leu Ser Tyr Asn Ser Gln Pro Phe Ser Met Lys
210 215 220

Gly Ile Gly His Asn Phe Ser Phe Val Thr His Leu Ser Met Leu Gln
225 230 235 240

Ser Leu Ser Leu Ala His Asn Asp Ile His Thr Arg Val Ser Ser His
245 250 255

Leu Asn Ser Asn Ser Val Arg Phe Leu Asp Phe Ser Gly Asn Gly Met
260 265 270

Gly Arg Met Trp Asp Glu Gly Gly Leu Tyr Leu His Phe Phe Gln Gly
275 280 285

Leu Ser Gly Val Leu Lys Leu Asp Leu Ser Gln Asn Asn Leu His Ile
290 295 300

Leu Arg Pro Gln Asn Leu Asp Asn Leu Pro Lys Ser Leu Lys Leu Leu
305 310 315 320

Ser Leu Arg Asp Asn Tyr Leu Ser Phe Phe Asn Trp Thr Ser Leu Ser
325 330 335

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Phe Leu Pro Asn Leu Glu Val Leu Asp Leu Ala Gly Asn Gln Leu Lys
            340                 345                 350

Ala Leu Thr Asn Gly Thr Leu Pro Asn Gly Thr Leu Leu Gln Lys Leu
        355                 360                 365

Asp Val Ser Ser Asn Ser Ile Val Ser Val Ala Pro Gly Phe Phe Ser
    370                 375                 380

Lys Ala Lys Glu Leu Arg Glu Leu Asn Leu Ser Ala Asn Ala Leu Lys
385                 390                 395                 400

Thr Val Asp His Ser Trp Phe Gly Pro Ile Val Met Asn Leu Thr Val
                405                 410                 415

Leu Asp Val Arg Ser Asn Pro Leu His Cys Ala Cys Gly Ala Ala Phe
            420                 425                 430

Val Asp Leu Leu Leu Glu Val Gln Thr Lys Val Pro Gly Leu Ala Asn
        435                 440                 445

Gly Val Lys Cys Gly Ser Pro Gly Gln Leu Gln Gly Arg Ser Ile Phe
    450                 455                 460

Ala Gln Asp Leu Arg Leu Cys Leu Asp Glu Val Leu Ser Trp Asp Cys
465                 470                 475                 480

Phe Gly Leu Ser Leu Leu Ala Val Ala Val Gly Met Val Val Pro Ile
                485                 490                 495

Leu His His Leu Cys Gly Trp Asp Val Trp Tyr Cys Phe His Leu Cys
            500                 505                 510

Leu Ala Trp Leu Pro Leu Leu Ala Arg Ser Arg Arg Ser Ala Gln Thr
        515                 520                 525

Leu Pro Tyr Asp Ala Phe Val Val Phe Asp Lys Ala Gln Ser Ala Val
    530                 535                 540

Ala Asp Trp Val Tyr Asn Glu Leu Arg Val Arg Leu Glu Glu Arg Arg
545                 550                 555                 560

Gly Arg Trp Ala Leu Arg Leu Cys Leu Glu Asp Arg Asp Trp Leu Pro
                565                 570                 575

Gly Gln Thr Leu Phe Glu Asn Leu Trp Ala Ser Ile Tyr Gly Ser Arg
            580                 585                 590

Lys Thr Leu Phe Val Leu Ala His Thr Asp Arg Val Ser Gly Leu Leu
        595                 600                 605

Arg Thr Ser Phe Leu Leu Ala Gln Gln Arg Leu Leu Glu Asp Arg Lys
    610                 615                 620

Asp Val Val Val Leu Val Ile Leu Arg Pro Asp Ala His Arg Ser Arg
625                 630                 635                 640
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Tyr Val Arg Leu Arg Gln Arg Leu Cys Arg Gln Ser Val Leu Phe Trp
645 650 655

Pro Gln Gln Pro Asn Gly Gln Gly Gly Phe Trp Ala Gln Leu Ser Thr
660 665 670

Ala Leu Thr Arg Asp Asn Arg His Phe Tyr Asn Gln Asn Phe Cys Arg
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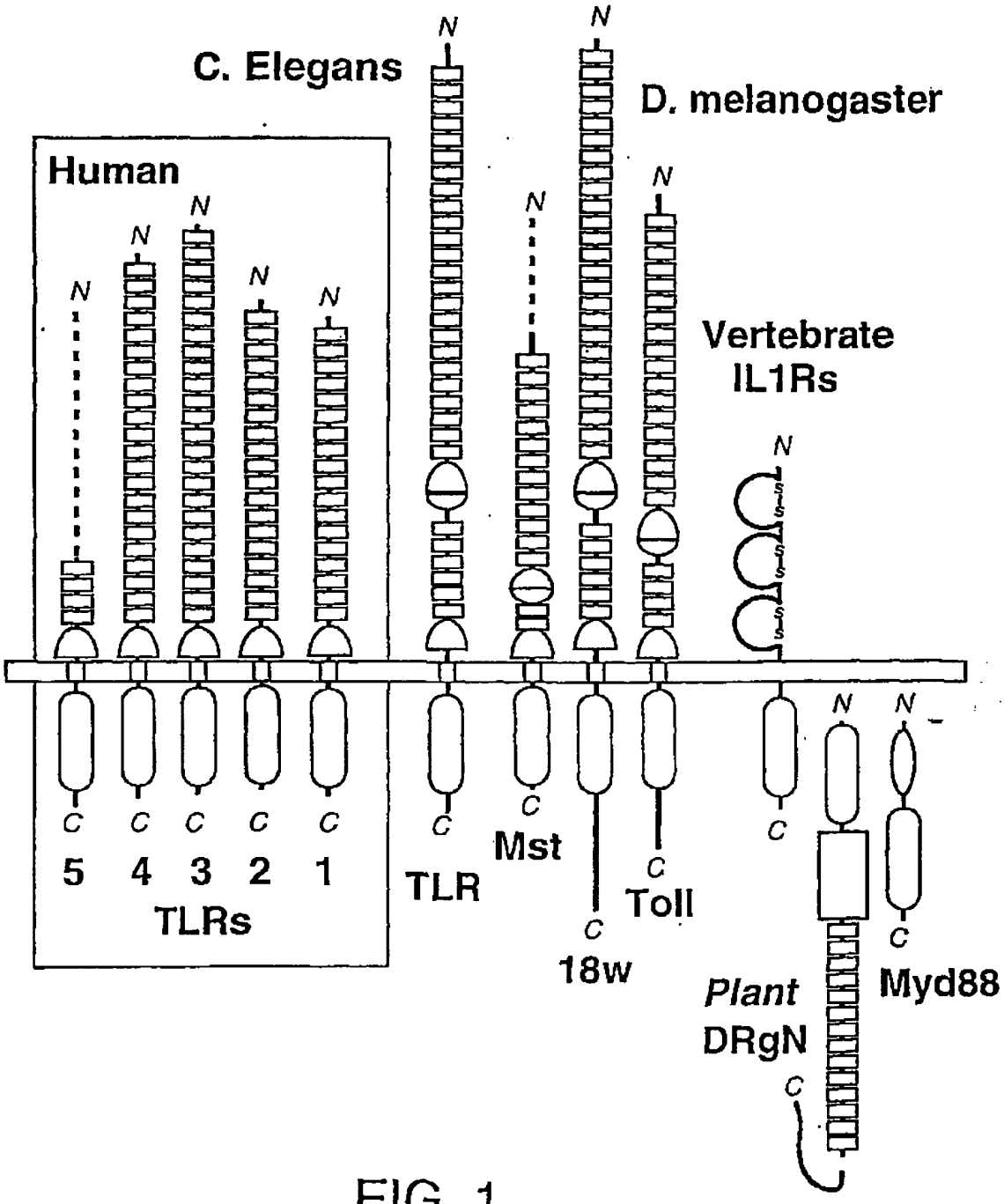
Gly Pro Thr Ala Glu
690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Toll
18w
Plant
DRgN
Myd88

FIG. 1

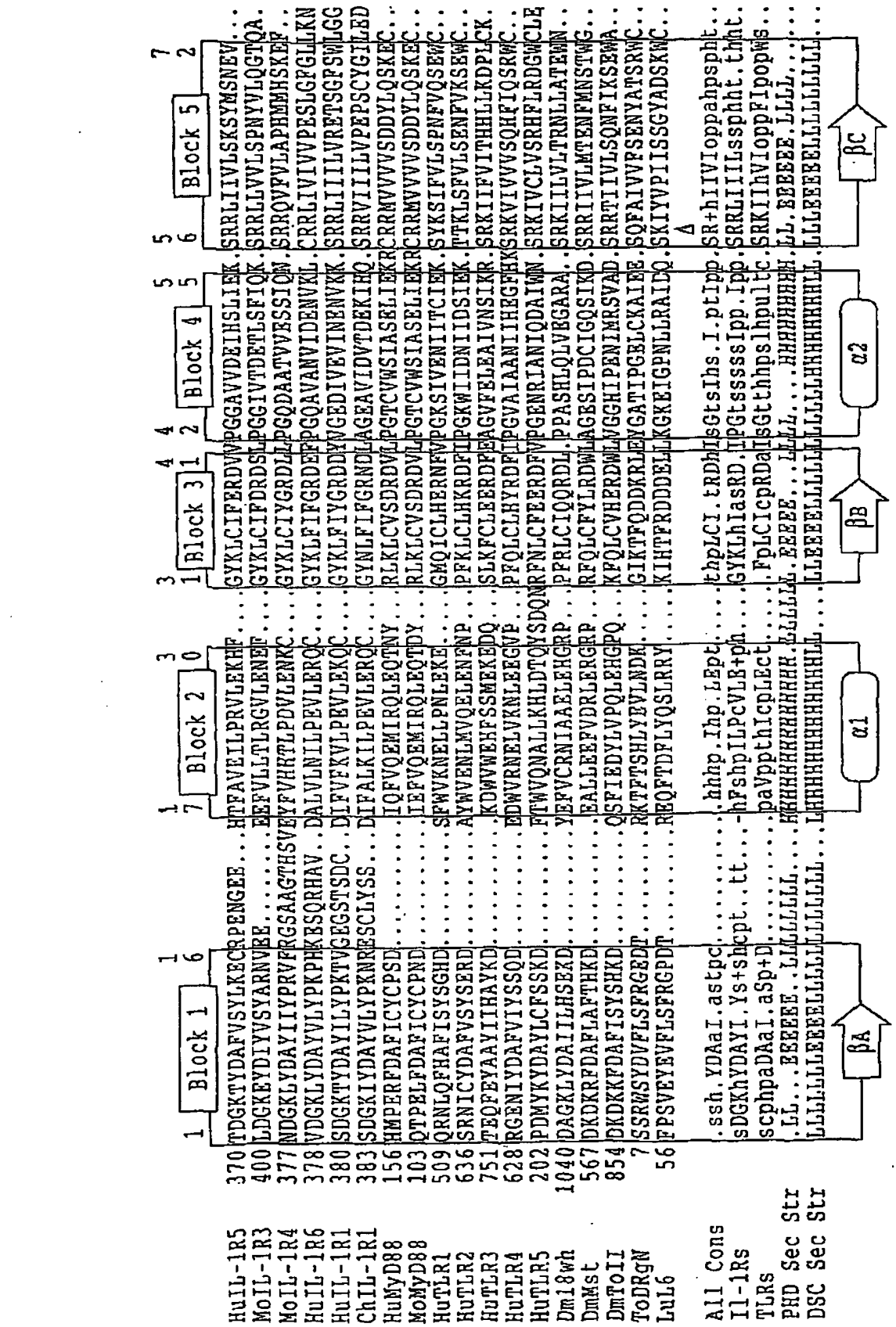
Block 1
Block 2
Block 3
Block 4
Block 5
βA
α1
βB
α2
βC
HuIL-1R5
MoIL-1R3
MoIL-1R4
HuIL-1R6
HuIL-1R1
ChIL-1R1
HuMyD88
MoMyD88
HuTLR1
HuTLR2
HuTLR3
HuTLR4
HuTLR5
Dm18wh
DmMst
DmToll
ToDRgN
LuL6
All Cons
Il-1Rs
TLRs
PHD Sec Str
DSC Sec Str

FIG. 2A

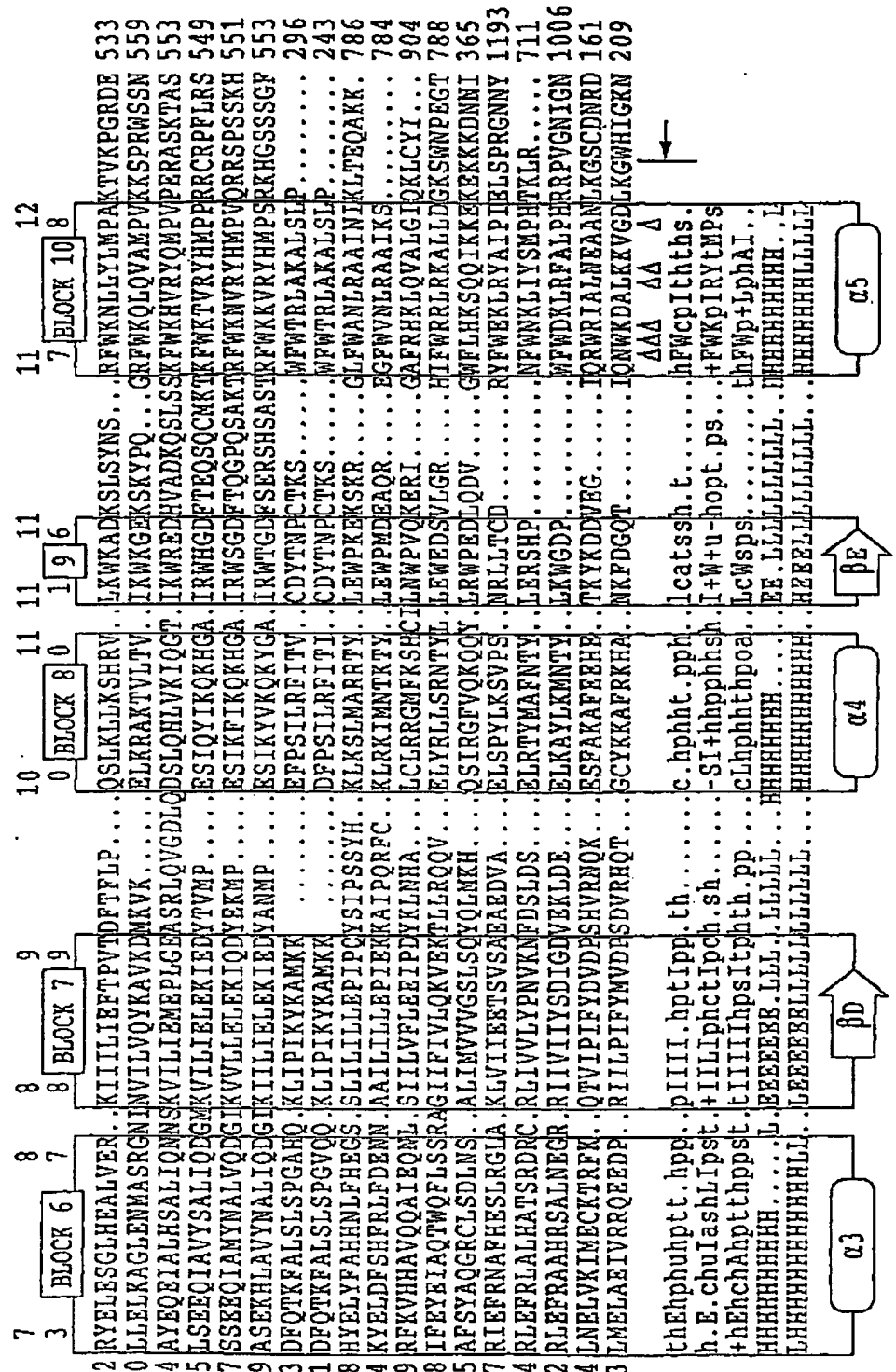

FIG. 2B

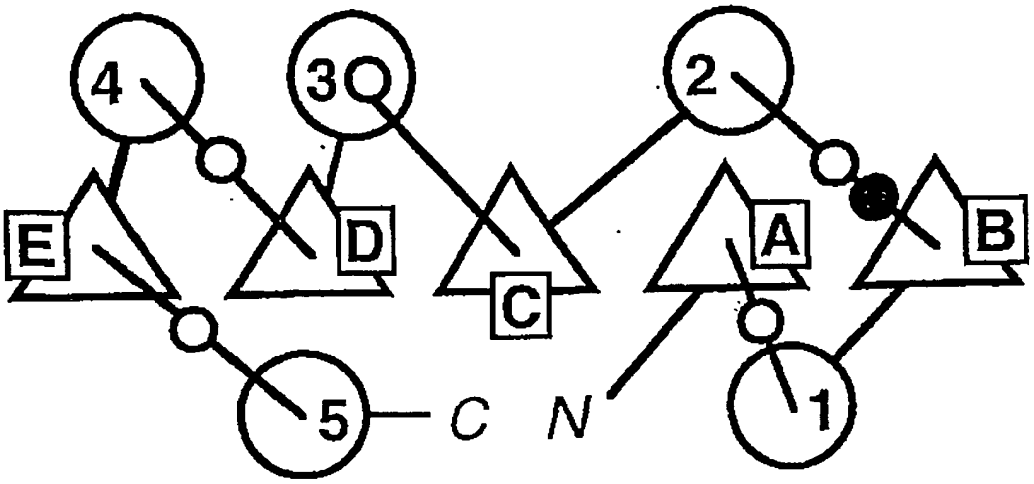
4
3
2
E
D
C
A
B
5
C
N
1

FIG. 2C

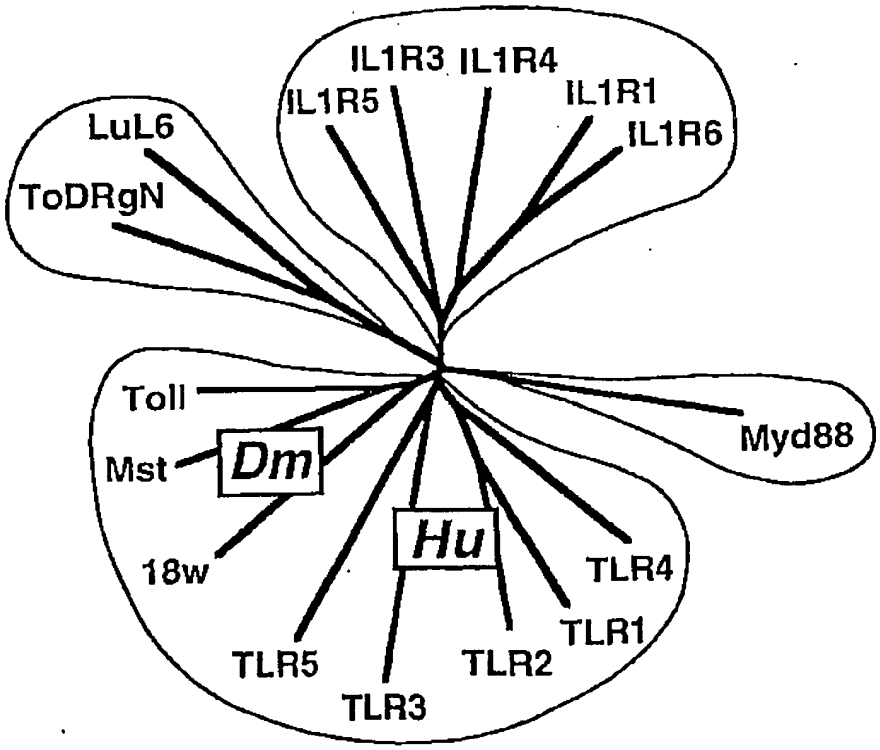
IL1R3
IL1R4
IL1R5
IL1R1
IL1R6
LuL6
ToDRgN
Toll
Myd88
Mst
Dm
Hu
18w
TLR4
TLR1
TLR5
TLR2
TLR3

FIG. 3

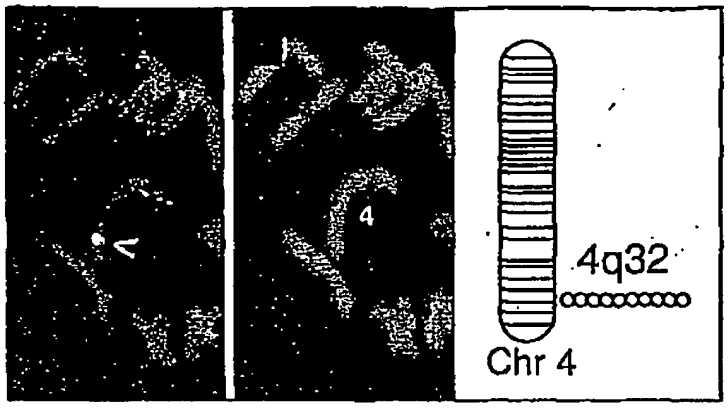

FIG. 4A

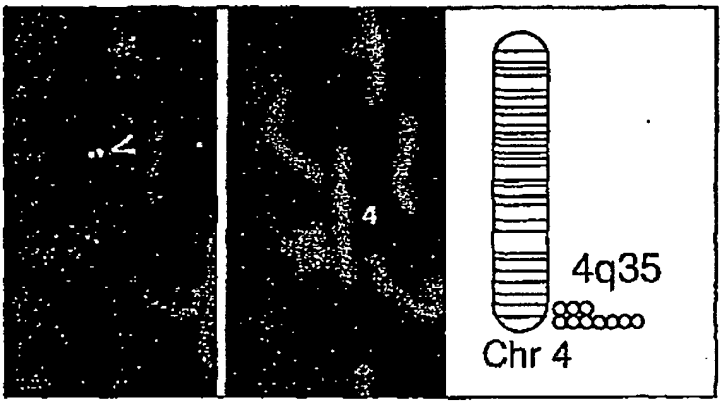

FIG. 4B

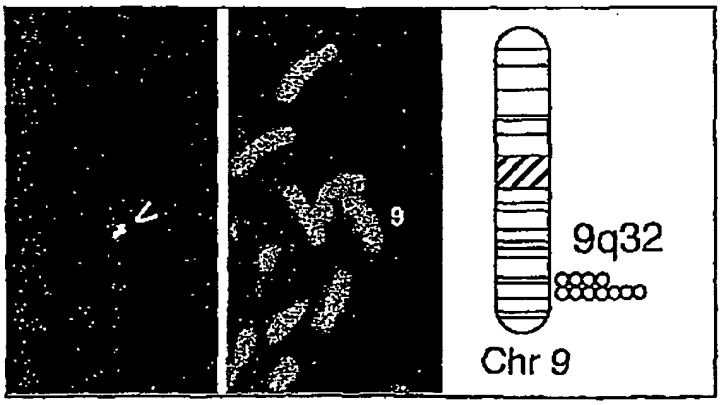

FIG. 4C

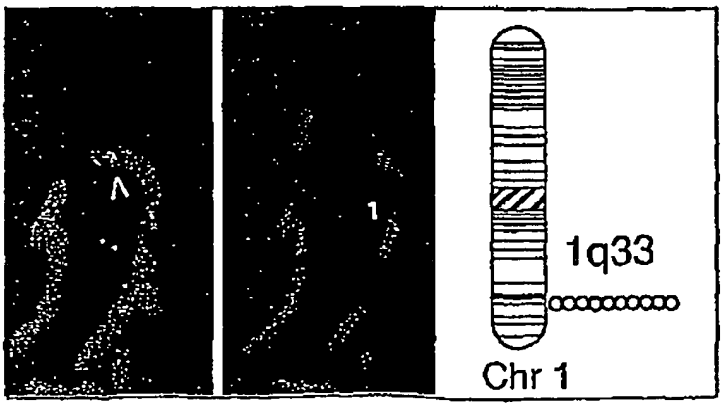

FIG. 4D

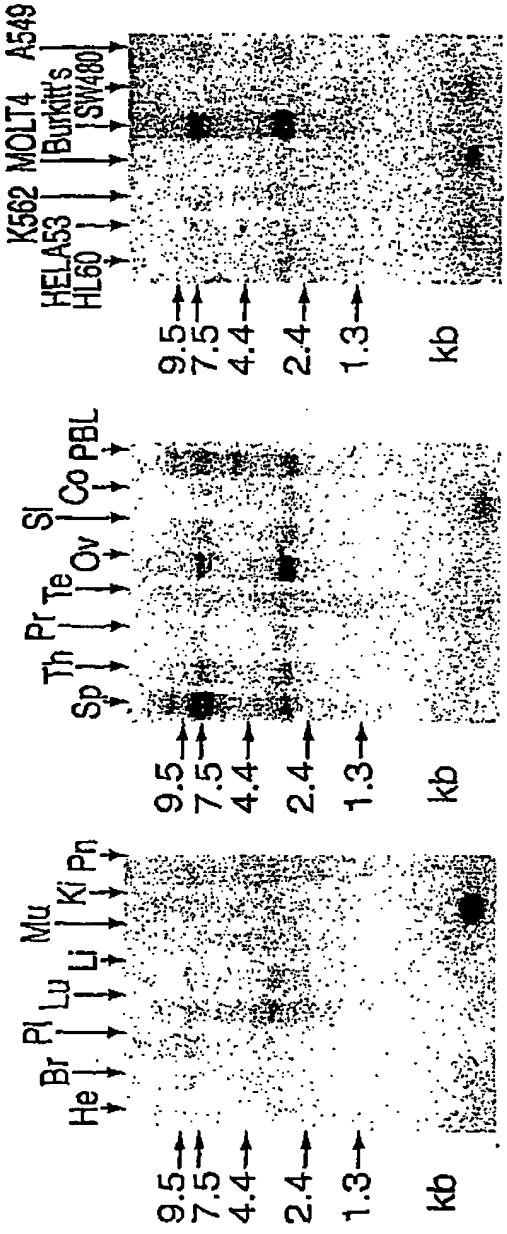

FIG. 5A

FIG. 5B

FIG. 5C

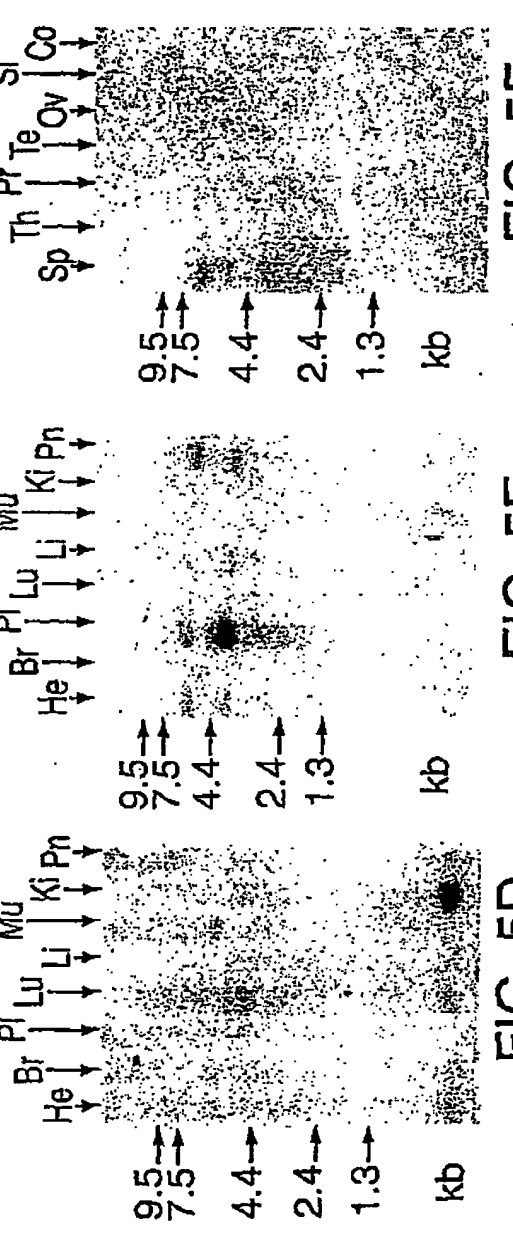

FIG. 5D

FIG. 5E

FIG. 5F

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

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(74) Agent: MCLAUGHLIN, Jaye, P.; Schering-Plough Corporation, Patent Dept. K-6-1 1990, 2000 Galloping Hill Road, Kenilworth, NJ 07033 (US).

(81) Designated States *(national)*: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, HR, HU, ID, IL, IN, IS, JP, KG, KR, KZ, LC, LK, LR, LT, LU, LV, MA, MD, MG, MK, MN, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, VN, YU, ZA.

(84) Designated States *(regional)*: ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

Declaration under Rule 4.17:
as to the applicant's entitlement to claim the priority of the earlier application (Rule 4.17(iii)) for all designations

Published:
— *with international search report*

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17 October 2002

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

WO 01/090151 A3

(54) Title: HUMAN RECEPTOR PROTEINS; RELATED REAGENTS AND METHODS

(57) Abstract: Nucleic acids encoding mammalian, e.g., human receptors, purified receptor proteins and fragments thereof. Antibodies, both polyclonal and monoclonal, are also provided. Methods of using the compositions for both diagnostic and therapeutic utilities are provided.

INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 01/16766

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 C07K14/705 C07K16/28 C12N15/12 C12N15/62 A61K38/17
A61K39/395 C12N5/10 C12N1/21

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 7 C12N C07K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)
EPO-Internal, BIOSIS, EMBL, WPI Data, PAJ, MEDLINE

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 98 50547 A (SCHERING CORP) 12 November 1998 (1998-11-12) DNA Seq ID no 3, Protein Seq ID no 4. ---	1-21
X	WO 99 20756 A (GENENTECH INC ;GODDARD AUDREY (US); GODOWSKI PAUL J (US); GURNEY A) 29 April 1999 (1999-04-29) DNA Seq ID no 11, Protein Seq ID no 12. --- -/--	1-21

[X] Further documents are listed in the continuation of box C.

[X] Patent family members are listed in annex.

* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
23 April 2002	26 07 2002

Name and mailing address of the ISA	Authorized officer
European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 351 epo nl, Fax: (+31-70) 340-3016	Aslund, J

Form PCT/ISA/210 (second sheet) (July 1992)

INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 01/16766

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	ROCK FERNANDO L ET AL: "A family of human receptors structurally related to Drosophila toll." PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES OF THE UNITED STATES, vol. 95, no. 2, 20 January 1998 (1998-01-20), pages 588-593, XP002197052 Jan. 20, 1998 ISSN: 0027-8424 Sequence of HuTLR2 in Figure 2. -& DATABASE EMBL [Online] EBI, Hinxton, UK; 3 October 1997 (1997-10-03) ROCK ET AL.: "Homo sapiens Toll-like receptor 2 (TLR2) mRNA, complete cds." Database accession no. U88878 XP002197054 the whole document ---	1-6, 11-18
X	CHAUDHARY PREET M ET AL: "Cloning and characterization of two toll/interleukin-1 receptor-like genes TIL3 and TIL4: Evidence for a multi-gene receptor family in humans." BLOOD, vol. 91, no. 11, 1 June 1998 (1998-06-01), pages 4020-4027, XP002197053 ISSN: 0006-4971 figure 1B -----	1-6, 11-18

Form PCT/ISA/210 (continuation of second sheet) (July 1992)

INTERNATIONAL SEARCH REPORT

PCT/US 01/16766

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

2. ☒ Claims Nos.: 19, 20
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
see FURTHER INFORMATION sheet PCT/ISA/210

3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

see additional sheet

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.

2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.

3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:

4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:
1-21 (all partially)

Remark on Protest

☐ The additional search fees were accompanied by the applicant's protest.

☐ No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.2

Claims Nos.: 19, 20

Present claims 20, 21 relate to a compound (ligand, agonist, antagonist) defined by reference to a desirable characteristic or property, namely the ability of modulating physiology or development of a DTLR protein. The claim cover all compounds having this characteristic or property, whereas the application provides support within the meaning of Article 6 PCT and/or disclosure within the meaning of Article 5 PCT only for antibodies. In the present case, the claims so lack support, and the application so lacks disclosure, that a meaningful search over the whole of the claimed scope is impossible. Independent of the above reasoning, the claims also lack clarity (Article 6 PCT). An attempt is made to define the compound by reference to a result to be achieved. Again, this lack of clarity in the present case is such as to render a meaningful search over the whole of the claimed scope impossible. Consequently, the search has been carried out for those parts of the claims which appear to be clear, supported and disclosed, namely those parts relating to an antibody with specificity for a DTLR protein.

The applicant's attention is drawn to the fact that claims, or parts of claims, relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

Invention 1: Clams 1-21 Partially

A DTLR2 protein of Seq Id 4.
Furthermore, nucleic acids and vectors encoding and expressing said polypeptide and derivatives such as fusion proteins, transformed host cells, antibodies, pharmaceutical compositions comprising said protein or antibodies directed to it.

Invention 2: Clams 1-21 Partially

A DTLR3 protein of Seq Id 6.
Furthermore, nucleic acids and vectors encoding and expressing said polypeptide and derivatives such as fusion proteins, transformed host cells, antibodies, pharmaceutical compositions comprising said protein or antibodies directed to it.

Invention 3: Clams 1-21 Partially

A DTLR4 protein of Seq Id 8.
Furthermore, nucleic acids and vectors encoding and expressing said polypeptide and derivatives such as fusion proteins, transformed host cells, antibodies, pharmaceutical compositions comprising said protein or antibodies directed to it.

Invention 4: Clams 1-21 Partially

A DTLR5 protein of Seq Id 10.
Furthermore, nucleic acids and vectors encoding and expressing said polypeptide and derivatives such as fusion proteins, transformed host cells, antibodies, pharmaceutical compositions comprising said protein or antibodies directed to it.

Invention 5: Clams 1-21 Partially

A DTLR6 protein of Seq Id 12, 28, 30.
Furthermore, nucleic acids and vectors encoding and expressing said polypeptide and derivatives such as fusion proteins, transformed host cells, antibodies, pharmaceutical compositions comprising said protein or antibodies directed to it.

Invention 6: Clams 1-21 Partially

A DTLR7 protein of Seq Id 16, 18, 37.
Furthermore, nucleic acids and vectors encoding and expressing said polypeptide and derivatives such as fusion proteins, transformed host cells, antibodies, pharmaceutical compositions comprising said protein or antibodies directed to it.

Invention 7: Clams 1-21 Partially

A DTLR8 protein of Seq Id 32, 39.
Furthermore, nucleic acids and vectors encoding and expressing said polypeptide and derivatives such as fusion proteins, transformed host cells, antibodies, pharmaceutical compositions comprising said protein or antibodies directed to it.

Invention 8: Clams 1-21 Partially

A DTLR9 protein of Seq Id 22, 41.
Furthermore, nucleic acids and vectors encoding and expressing said polypeptide and derivatives such as fusion proteins, transformed host cells, antibodies, pharmaceutical compositions comprising said protein or antibodies directed to it.

Invention 9, Claims 1-21 Partially

A DTLR10 protein of Seq Id 34, 43, 45.
Furthermore, nucleic acids and vectors encoding and expressing said polypeptide and derivatives such as fusion proteins, transformed host cells, antibodies, pharmaceutical compositions comprising said protein or antibodies directed to it.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No
PCT/US 01/16766

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9850547 A	12-11-1998	AU 740333 B2	01-11-2001
		AU 7175498 A	27-11-1998
		BR 9808747 A	11-07-2000
		CN 1263555 T	16-08-2000
		EP 0980429 A2	23-02-2000
		JP 2002514083 T	14-05-2002
		NO 995458 A	08-11-1999
		NZ 338073 A	28-09-2001
		PL 336635 A1	03-07-2000
		SK 146599 A3	11-07-2000
		WO 9850547 A2	12-11-1998
		HU 0001462 A2	28-07-2000
WO 9920756 A	29-04-1999	AU 1070399 A	10-05-1999
		CA 2305385 A1	29-04-1999
		EP 1025227 A2	09-08-2000
		WO 9920756 A2	29-04-1999
		US 2002086368 A1	04-07-2002
		US 2002072496 A1	13-06-2002
		US 2002072067 A1	13-06-2002
		US 2002072092 A1	13-06-2002
		US 2002072497 A1	13-06-2002
		ZA 9809147 A	07-04-2000
		US 2002052019 A1	02-05-2002
		WO 9946281 A2	16-09-1999
		CA 2328895 A1	09-12-1999
		WO 9963088 A2	09-12-1999